



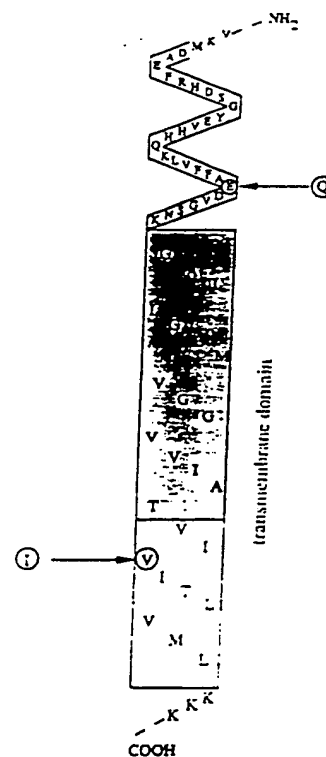
## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/GB92/00123 <b>(22) International Filing Date:</b> 21 January 1992 (21.01.92) <b>(30) Priority data:</b> 9101307.8 21 January 1991 (21.01.91) GB 9118445.7 28 August 1991 (28.08.91) GB <b>(71) Applicant (for all designated States except US):</b> IMPERIAL COLLEGE OF SCIENCE, TECHNOLOGY & MEDICINE[GB/GB]; Sherfield Building, Exhibition Road, London SW7 2AZ (GB). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only) :</b> HARDY, John, Anthony [GB/GB]; 187 Drakefell Road, Brockley, London SE4 (GB). CHARTIER-HARLIN, Marie-Christine [FR/GB]; 63 Francis Road, Leyton, London E10 6PN (GB). GOATE, Alison, Mary [GB/GB]; 100 High Street, Hampton Wick, Kingston, Surrey KT1 4DQ (GB). OWEN, Michael, John [GB/GB]; Four Hedges, Castlehill, Llanblethian, Nr. Cowbridge, South Glamorgan CF7 7JB (GB). MULLAN, Michael, John [GB/US]; Suncoast Gerontology Centre, University of South Florida, 12901 Bruce B. Downs Blvd. MDC 50, Tampa, FL 33612 (US).		<b>(74) Agents:</b> SHEARD, Andrew, Gregory et al.; Kilburn & Strode, 30 John Street, London WC1N 2DD (GB). <b>(81) Designated States:</b> AT (European patent), AU, BB, BE (European patent), BF (OAPI patent), BG, BJ (OAPI patent), BR, CA, CF (OAPI patent), CG (OAPI patent), CH (European patent), CI (OAPI patent), CM (OAPI patent), CS, DE (European patent), DK (European patent), ES (European patent), FI, FR (European patent), GA (OAPI patent), GB (European patent), GN (OAPI patent), GR (European patent), HU, IT (European patent), JP, KP, KR, LK, LU (European patent), MC (European patent), MG, ML (OAPI patent), MN, MR (OAPI patent), MW, NL (European patent), NO, PL, RO, RU, SD, SE (European patent), SN (OAPI patent), TD (OAPI patent), TG (OAPI patent), US. <b>Published</b> With international search report.

(54) Title: TEST AND MODEL FOR ALZHEIMER'S DISEASE

## (57) Abstract

Model systems of Alzheimer's disease comprise a DNA sequence encoding an amyloid precursor protein (APP) isoform or fragment that has an amino acid substitution. The substituted amino acid may be other than valine at the amino acid position corresponding to amino acid residue position 717 of APP770. Methods of determining genetic predisposition to Alzheimer's disease are also disclosed.

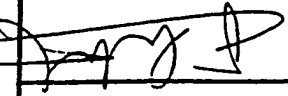


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## TEST AND MODEL FOR ALZHEIMER'S DISEASE

### BACKGROUND OF THE INVENTION

Alzheimer's disease is a progressive disease known generally as senile dementia. Broadly speaking the disease falls into two categories, namely late onset and early onset. Late onset, which occurs in old age (65 + years), may be caused by the natural atrophy of the brain occurring at a faster rate and to a more severe degree than normal. Early onset Alzheimer's disease is much more infrequent but shows a pathologically identical dementia with diffuse brain atrophy which develops well before the senile period, *i.e.*, between the ages of 35 and 60 years. There is evidence that one form of this type of Alzheimer's disease shows a tendency to run in families and is therefore known as familial Alzheimer's disease (FAD).

In both types of Alzheimer's disease the pathology is the same but the abnormalities tend to be more severe and more widespread in cases beginning at an earlier age. The disease is characterized by two types of lesions in the brain, these are senile plaques and neurofibrillary tangles.

Senile plaques are areas of disorganized neuropil up to 150 $\mu$ m across with extracellular amyloid deposits at the center. Neurofibrillary tangles are intracellular deposits of amyloid protein consisting of two filaments twisted about each other in pairs.

The major protein subunit,  $\beta$ -amyloid protein, of the amyloid filaments of the senile plaque is a highly aggregating small polypeptide of approximate relative molecular mass 4,500. This protein is a cleavage product of a much larger precursor protein called amyloid precursor protein (APP).

At present there is no known effective therapy for the various forms of Alzheimer's disease (AD). However, there are several other forms of dementia for which treatment is available and which give rise to progressive intellectual deterioration closely resembling the dementia associated with Alzheimer's disease.

A diagnostic test for AD would therefore provide a valuable tool in the diagnosis and treatment of these other conditions, by way of being able to exclude Alzheimer's disease. It will also be of value when a suitable therapy does become available.

5                   Also important is the development of experimental models of Alzheimer's disease that can be used to define further the underlying biochemical events involved in AD pathogenesis. Such models could presumably be employed, in one application, to screen for agents that alter the degenerative course of Alzheimer's disease. For example, a model system of Alzheimer's disease could  
10 be used to screen for environmental factors that induce or accelerate the pathogenesis of AD. In contradistinction, an experimental model could be used to screen for agents that inhibit, prevent, or reverse the progression of AD. Presumably, such models could be employed to develop pharmaceuticals that are effective in preventing, arresting, or reversing AD.

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#### SUMMARY OF THE INVENTION

The present invention provides model systems of Alzheimer's disease, wherein the model system comprises a DNA sequence encoding an amyloid precursor protein (APP) isoform or fragment that has an amino acid other  
20 than valine at the amino acid position corresponding to amino acid residue position 717 of APP770.

In a first embodiment, the present invention provides an isolated DNA sequence that encodes an amyloid precursor protein (APP) isoform or fragment that has an amino acid other than valine at the amino acid position  
25 corresponding to amino acid residue position 717 of APP770.

In a second embodiment, the present invention provides a transgenic nonhuman animal that harbors at least one integrated copy of a human DNA sequence that encodes an amyloid precursor protein (APP) isoform or fragment that has an amino acid other than valine at the amino acid position corresponding

to amino acid residue position 717 of APP770.

In a third embodiment, the present invention provides a transgenic nonhuman animal wherein at least one of the endogenous nonhuman APP alleles has been completely or partially replaced by all or a portion of a human APP gene that includes a codon 717 that does not encode valine.

In a fourth embodiment, the present invention provides cells, typically mammalian cells and preferably mammalian cells of the neural, glial, or astrocytic lineage, that have been transformed or transfected with a heterologous DNA sequence, or have been derived from a transgenic nonhuman animal, wherein the cells express an amyloid precursor protein (APP) isoform or fragment that has an amino acid other than valine at the amino acid position corresponding to amino acid residue position 717 of APP770. In accordance with standard protocols, cultured human cells, either primary cultures or immortalized cell lines, may be transfected, either transiently or stably, with a mutant APP allele so that the cultured human cell expresses a mutant APP polypeptide.

In a fifth embodiment, the present invention provides a method of producing transgenic nonhuman animals and transformed cells that contain a DNA sequence encoding an amyloid precursor protein (APP) isoform or fragment that has an amino acid other than valine at the amino acid position corresponding to amino acid residue position 717 of APP770.

In a sixth embodiment, the present invention provides a method of producing, free from other human proteins, a human amyloid precursor protein (APP) isoform or fragment that has an amino acid other than valine at the amino acid position corresponding to amino acid residue position 717 of APP770.

In a seventh embodiment, the present invention provides a human amyloid precursor protein (APP) isoform or fragment, free from other human proteins, that has an amino acid other than valine at the amino acid position corresponding to amino acid residue position 717 of APP770.

In an eighth embodiment, the invention provides a method for detecting an APP allele that is linked (i.e., cosegregates with) a genetic predisposition to Alzheimer's disease, particularly early onset AD, wherein such a pathognomonic APP allele is detected by determining that codon 717 of the allele  
5 does not encode valine. Preferably, a pathognomonic APP allele is detected when codon 717 is determined to encode either isoleucine, glycine, or phenylalanine. Thus, methods for locating the presence of genetic alterations associated with Alzheimer's disease are provided. This diagnostic method may be used to predict the development of the disease prior to onset, for genetic screening, or to detect  
10 a specific mutation in an experimental nonhuman animal or a cell.

In a ninth embodiment, the invention provides a human variant APP polypeptide free of other human proteins, typically present in a cell of a nonhuman animal. The invention also relates to an isolated nucleic acid encoding such a polypeptide and to uses and applications of such nucleic acid as are described  
15 above in relation to the specific embodiment of the invention which involves an amino acid substitution at position 717 (as defined in relation to APP770).

According to one aspect of the invention there is provided a method for detecting the presence, in a nucleic acid or other sample removed from a subject, of the gene for Alzheimer's disease comprising identifying a genetic  
20 alteration in a gene sequence coding for APP. Such genetic alterations may include mutations, insertions or deletions.

#### BRIEF DESCRIPTION OF THE FIGURES

Fig. 1 illustrates a first pedigree in which early onset AD is  
25 apparently inherited as an autosomal dominant disorder. The average age of onset in this family is  $57 \pm 5$  years. Black symbols denote affected individuals and oblique lines indicate individuals who are deceased. Females are denoted by circles and males by squares. Triangles are used in the present generation to preserve anonymity. In generation II the spouses of the two affected brothers were



sisters. Samples were available from the 13 individuals whose haplotypes are illustrated, from a further 19 children and spouses of these individuals and from 7 more distantly related unaffected individuals. Beneath the pedigree are ideograms of the two chromosomes 21 in each individual of the third generation at four loci on the long arm of the chromosome. The linkage data suggest that the black chromosomes were inherited from the affected fathers.

Fig. 2 shows an autoradiograph of a sequencing gel from part of exon 17 of the APP gene in a normal and an affected individual from the Fig. 1 pedigree showing a single base pair change at base pair 2149 in the affected individual. This C to T transition leads to an amino acid substitution of a valine by an isoleucine at codon 717.

Fig. 3 shows part of the amino acid sequence encoded by exons 16 and 17 of the APP gene showing the mutation valine to isoleucine (V to I) within the transmembrane domain and the mutation causing HCHWA-D (E to Q) in the extracellular domain. The shaded region of the transmembrane domain and the boxed amino acids of the extracellular domain represent the sequence of the deposited  $\beta$ -amyloid peptide. Adapted from Kang *et al.* (1987) *Nature* 325:733.

Fig. 4 shows *Bcl*I digests of the exon 17 PCR product from unaffected and affected individuals in an early onset AD family showing co-segregation of the restriction site and the disease.

Fig. 5 shows the pedigree of family F19, together with D21S210 data.

Fig. 6 shows APP exon 17 sequences in an affected and unaffected member of F19. In the affected member there is a G- > T transition at position 2150.

Fig. 7 shows the sequence of APP695.

Fig. 8 shows the sequence of APP751.

Fig. 9 shows the sequence of APP770.

## DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

The accumulation of  $\beta$ -amyloid protein (A4) in particular brain regions is one of the main pathologic characteristics of Alzheimer's disease. The  $\beta$ -amyloid protein is an approximately 4 kD protein (39 to 42 amino acids) which is derived, as an internal cleavage product, from one or more isoforms of a larger amyloid precursor protein (APP). There are at least five distinct isoforms of APP containing 563, 695, 714, 751, and 770 amino acids, respectively (Wirak et al. (1991) *Science* 253:323). These isoforms of APP are generated by alternative splicing of primary transcripts of a single gene, designated the APP gene, which is located on human chromosome 21. It is known that most of the APP isoforms are glycosylated transmembrane proteins (Goldgaber et al. (1987) *Science* 235:877), and that four of the isoforms, AA563, APP714, APP751 and APP770, have a protease inhibitor domain that is homologous to the Kunitz type of serine protease inhibitors. The  $\beta$ -amyloid (A4) segment comprises approximately half of the transmembrane domain and approximately the first 28 amino acids of the extracellular domain of an APP isoform.

Proteolytic processing of APP *in vivo* is a normal physiological process. Carboxy-terminal truncated forms of APP695, APP751, and APP770 are present in brain and cerebrospinal fluid (Palmert et al. (1989) *Proc. Natl. Acad. Sci. U.S.A.* 86:6338; Weidemann et al. (1989) *Cell* 57:115) and result from cleavage of the APP isoform at a constitutive cleavage site within the A4 peptide domain of an APP isoform (Esch et al. (1990) *Science* 248:1122). Normal proteolytic cleavage at the constitutive cleavage site yields a large (approximately 100 kD) soluble, N-terminal fragment that contains the protease inhibitor domain in some isoforms, and a 9-10 kD membrane-bound, C-terminal fragment that includes most of the A4 domain.

Generation of pathogenic  $\beta$ -amyloid (A4) protein appears to be the result of aberrant or alternative proteolytic processing of APP, such that normal cleavage at the constitutive site within the A4 domain does not occur, but rather

cleavage occurs at two specific sites which flank the A4 domain. One of these aberrant cleavage sites is in the transmembrane domain and the other aberrant cleavage site is located approximately at the N-terminus of the first 28 amino acids of the extracellular domain (see Fig. 3). Such aberrant proteolytic cleavage produces the  $\beta$ -amyloid A4 polypeptide which is prone to forming dense amyloidogenic aggregates that are resistant to proteolytic degradation and removal. The resultant  $\beta$ -amyloid aggregates presumably are involved in the formation of the abundant amyloid plaques and cerebrovascular amyloid that are the neuropathological hallmarks of Alzheimer's disease. However, the exact aberrant cleavage sites are not always precise;  $\beta$ -amyloid molecules isolated from the brain of a patient with AD show N- and C- terminal heterogeneity. Therefore, the aberrant cleavage pathway may involve either sequence-specific proteolysis followed by exopeptidase activity (creating end-heterogeneity), or may not be sequence-specific.

The APP gene is known to be located on human chromosome 21. A locus segregating with familial Alzheimer's disease has been mapped to chromosome 21 (Hyslop *et al.* (1987) *Science* 235:885) close to the APP gene. Recombinants between the APP gene and the AD locus have been previously reported (Schellenberg *et al.* (1988) *Science* 241:1507). The data appeared to exclude the APP gene as the site of any mutation that might cause FAD (Van Broekhoven *et al.* (1987) *Nature* 329:153; Tanzi *et al.* (1987) *Nature* 329:156).

Recombinant DNA technology provides several techniques for analyzing genes to locate possible mutations. For example, the polymerase chain reaction (Bell (1989) *Immunology Today*, 10:351) may be used to amplify specific sequences using intronic primers, which can then be analyzed by direct sequencing.

Researchers working in the area of the hereditary cerebral haemorrhage with amyloidosis of the Dutch type ("HCHWA-D") (Levy *et al.* (1990) *Science* 248:11224) found a substitution of Glu to Gln at residue 618 (using

the APP695 numbering system) in APP which is thought to result in the deposition of  $\beta$ -amyloid in the cerebral vessels of these patients. The present inventors have identified a single base substitution, a C to T transition at base pair 2149, has been found in part of the sequence of the APP gene in some cases of familial Alzheimer's disease. This base pair transition leads to an amino acid substitution, i.e., valine to isoleucine at amino acid 717 (APP<sub>770</sub>) (see Yoshikai *et al.* (1990) Gene 87:257), close to the C-terminus of the  $\beta$ -amyloid protein. This suggests that some cases of Alzheimer's disease are caused by mutations in the APP gene, and specifically mutations that change codon 717 such that it encodes an amino acid other than valine.

Additionally, a further single base substitution, a T to G transition at adjacent base pair 2150, has been found in part of the sequence of the APP gene in other cases of familial Alzheimer's disease. This base pair transition leads to a different amino acid substitution, namely valine to glycine, at amino acid 717, thereby strengthening the argument that some cases of Alzheimer's disease are caused by mutations in the APP gene, specifically at codon 717.

It is now clear that a mutation in the APP gene locus that results in a substitution of isoleucine for valine at codon 717 (residue 642 in APP695) gives rise to AD in some families (Goate *et al.* (1991) Nature 349:704). A second APP allelic variant wherein glycine is substituted for valine at codon 717 is now identified, and is so closely linked to the AD phenotype as to indicate that allelic variants at codon 717 of the APP gene, particularly those encoding an amino acid other than valine, and more particularly those encoding a isoleucine, glycine, or phenylalanine, are pathogenic and/or pathognomonic alleles (Chartier-Harlin *et al.* (1991) Nature 353:844).

Proteolysis on either side of the  $\beta$ -amyloid (A4) region of APP may be enhanced or qualitatively altered by the specific mutations at codon 717, increasing the rate of  $\beta$ -amyloid deposition and aggregation. Such codon 717 mutations may increase  $\beta$ -amyloid formation by providing a poorer substrate for

the main proteolytic pathway (cleavage at the constitutive site) or a better substrate for a competing, alternative cleavage pathway (at aberrant cleavage sites).

### DEFINITIONS

5                   A number of terms and expressions are used throughout the specification and, to facilitate the understanding thereof, the following definitions are provided:

                  As used herein, "exon" refers to any segment of an interrupted gene that is represented in the mature RNA product.

10                   As used herein, "intron" refers to a segment of an interrupted gene that is not represented in the mature RNA product. Introns are part of the primary nuclear transcript but are spliced out to produce mRNA, which is then transported to the cytoplasm.

                  As used herein, the phrase "gene sequence coding for amyloid protein precursor" may be interpreted to mean the DNA and cDNA sequence as detailed by Yoshikai *et al.* (1990) Gene 87:257 and Kang *et al.*, loc. cit., together with the promoter DNA sequence as described by Salbaum *et al.* (1988) EMBO 7(9):2807.

20                   As used herein, the terms "label" or "labeled" refers to incorporation of a detectable marker (e.g., by incorporation of a radiolabeled nucleotide or by end-labeling with a terminal radiolabeled phosphate). DNA or RNA is typically labeled by incorporation of a radiolabeled nucleotide ( $H^3$ ,  $C^{14}$ ,  $S^{35}$ ,  $P^{32}$ ) or a biotinylated nucleotide that can be detected by marked avidin (e.g., avidin containing a fluorescent marker or enzymatic activity) or digoxigeninylated  
25                   nucleotide that can be detected by marked specific antibody.

                  As used herein, "isoform", "APP", and "APP isoform" refer to a polypeptide that is encoded by at least one exon of the APP gene (Kitaguchi *et al.* (1988) Nature 331:530; Ponte *et al.*, ibid., p.525; R.E. Tanzi, ibid., p.528; de Sauvage and Octave (1989) Science 245:651; Golde *et al.* (1990) Neuron 4:253).

An APP isoform may be encoded by an APP allele (or exon thereof) that is associated with a form of Alzheimer's disease or that is not associated with an AD disease phenotype.

5 The term " $\beta$ -amyloid gene" is used herein as a synonym for the APP gene, as  $\beta$ -amyloid is a protein product produced by a post-translational cleavage of an APP gene product.

As used herein, "fragment" refers to a polypeptide of at least about 9 amino acids, typically 50 to 75, or more, wherein the polypeptide contains an amino acid core sequence (listed in order from amino- to carboxy-terminal direction):

-Ile-Ala-Thr-Val-Ile-X-Ile-Thr-Leu- [SEQ ID NO:6]

where X is any of the twenty conventional amino acids except valine, and particularly where X is isoleucine, glycine, or phenylalanine. A fragment may be a truncated APP isoform, modified APP isoform (as by amino acid substitutions, deletions, or additions outside of the core sequence), or other variant polypeptide sequence, but is not a naturally-occurring APP isoform or  $\beta$ -amyloid polypeptide that is present in a human individual, whether affected by AD or not. If desired, the fragment may be fused at either terminus to additional amino acids, which may number from 1 to 20, typically 50 to 100, but up to 250 to 500 or more.

20 As used herein, "APP751" and "APP770" refer, respectively, to the 751 and 770 amino acid residue long polypeptides encoded by the human APP gene (Ponte et al. loc. cit.; Kitaguchi et al. loc. cit.; Tanzi et al. loc. cit.).

As used herein, "codon 717" refers to the codon (i.e., the trinucleotide sequence) that encodes the 717th amino acid position in APP770, or the amino acid position in an APP isoform or fragment that corresponds to the 717th position in APP770. For example but not limitation, a 670 residue long fragment that is produced by truncating APP770 by removing the 100 N-terminal amino acids has its 617th amino acid position corresponding to codon 717. In fact, as used herein, codon 717 refers to the codon that encodes the 698th amino acid

residue of APP751 [SEQ ID NO:2] and the 642nd amino acid residue of APP695 [SEQ ID NO:1].

As used herein, "human APP isoform or fragment" refers to an APP isoform or fragment that contains a sequence of at least 9 consecutive amino acids that is identical to a sequence in a human APP770, APP751, or APP695 protein that occurs naturally in a human individual, and wherein an identical sequence is not present in an APP protein that occurs naturally in a nonhuman species.

A nucleic acid is "operably linked" when it is placed into a functional relationship with another nucleic acid sequence. For instance, a promoter or enhancer is operably linked to a coding sequence if it affects the transcription of the sequence. With respect to transcription regulatory sequences, operably linked means that the DNA sequences being linked are contiguous and, where necessary to join two protein coding regions, contiguous and in reading frame.

The term "corresponds to" is used herein to mean that a sequence is homologous (i.e., is identical, not strictly evolutionarily related) to all or a portion of a reference sequence. In contradistinction, the term "complementary to" is used herein to mean that the complementary sequence is homologous to all or a portion of a reference sequence. For illustration, the nucleotide sequence "TATAC" corresponds to a reference sequence "TATAC" and is complementary to a reference sequence "GTATA".

The term "transcriptional enhancement" is used herein to refer to functional property of producing an increase in the rate of transcription of linked sequences that contain a functional promoter.

The term "agent" is used herein to denote a chemical compound, a mixture of chemical compounds, a biological macromolecule, or an extract made from biological materials such as bacteria, plants, fungi, or animal (particularly mammalian) cells or tissues. Agents are evaluated for potential biological activity by inclusion in screening assays described hereinbelow.

As used herein, the term "mutant" refers to APP alleles having missense mutations that are pathognomonic for a genetic predisposition for developing AD; specifically a mutation at codon 717 (as referenced by the amino acid sequence in APP770) of the APP gene, such that codon 717 encodes one of the nineteen amino acids that are not valine (i.e., glycine, methionine, alanine, serine, isoleucine, leucine, threonine, proline, histidine, cysteine, tyrosine, phenylalanine, glutamic acid, tryptophan, arginine, aspartic acid, asparagine, lysine, and glutamine), but preferably isoleucine, glycine, or phenylalanine. Thus a mutant APP770 polypeptide is an APP770 polypeptide that has an amino acid residue at position 717 that is not valine. Other mutant APP isoforms comprise a non-valine amino acid at the amino acid residue position that corresponds to codon 717 (i.e., that is encoded by codon 717). Similarly, a mutant APP allele or a variant APP codon 717 allele is an APP allele that encodes an amino acid other than valine at codon 717 (referenced to the human APP770 deduced translation as described in the "codon 717" definition, *supra*), preferably isoleucine, glycine, or phenylalanine. Hence, an APP allele that encodes valine at codon 717 is a "wild-type" APP allele.

It is apparent to one of skill in the art that nucleotide substitutions, deletions, and additions may be incorporated into the polynucleotides of the invention. However, such nucleotide substitutions, deletions, and additions should not substantially disrupt the ability of the polynucleotide to hybridize to one of the polynucleotide sequences shown in Figs. 5 and 6 under hybridization conditions that are sufficiently stringent to result in specific hybridization.

"Specific hybridization" is defined herein as the formation of hybrids between a probe polynucleotide (e.g., a polynucleotide of the invention which may include substitutions, deletion, and/or additions) and a specific target polynucleotide (e.g., a polynucleotide having the sequence ), wherein the probe preferentially hybridizes to the specific target such that, for example, a band corresponding to a variant APP allele or restriction fragment thereof, can be



identified on a Southern blot, whereas a corresponding wild-type APP allele (i.e., one that encodes valine at codon 717) is not identified or can be discriminated from a variant APP allele on the basis of signal intensity. Hybridization probes capable of specific hybridization to detect a single-base mismatch may be designed according to methods known in the art and described in Maniatis et al., Molecular Cloning: A Laboratory Manual, 2nd Ed., (1989), Cold Spring Harbor, N.Y. and Berger and Kimmel, Methods in Enzymology, Volume 152, Guide to Molecular Cloning Techniques (1987), Academic Press, Inc., San Diego, CA; Gibbs et al. (1989) Nucleic Acids Res. 17:2437; Kwok et al. (1990) Nucleic Acids Res. 18:999; Miyada et al. (1987) Methods Enzymol. 154:94, each of which is incorporated herein by reference. The  $T_m$  for oligonucleotides is calculated under standard conditions (1 M NaCl) to be  $[4^{\circ}\text{C} \times (\text{G}+\text{C}) + 2^{\circ}\text{C} \times (\text{A}+\text{T})]$ . While the conditions of PCR differ from the standard conditions, this  $T_m$  is used as a guide for the expected relative stabilities of oligonucleotides. Allele-specific primers are typically 13-15 nucleotides long, sometimes 16-21 nucleotides long, or longer; when short primers are used, such as a 14 nucleotide long primer, low annealing temperatures are used, typically 44 to 50°C, occasionally somewhat higher or lower depending on the base composition of the sequence(s).

## DESCRIPTION OF THE PREFERRED EMBODIMENTS

### Detection of Mutant Codon 717 APP Alleles

In an embodiment of the invention, the method involves identifying a genetic alteration at amino acid 717, which may cause the consensus Val to be changed, for example, to another hydrophobic residue. This will generally be performed on a specimen removed from the subject. Hydrophobic residues include Leu, Ala, Ile and Gly, the first three of which have aliphatic side chains. Phe also has a hydrophobic residue which may be appropriate. As indicated above, preferred residues include Ile, Gly, and Phe (Murrell *et al*, (1991) *Science* 254:97).

The fact that these mutations discussed above are at the same codon may be a coincidence, but this seems unlikely on statistical grounds. There are two possibilities that may explain these data. First, substitution of the valine residue at codon 717 may result in increased beta-amyloid deposition due to changes in APP metabolism. Secondly, the variation in the sequence around this position may result in increased translation of APP mRNAs and thus cause AD by a route analogous to that by which AD is believed to be caused in Down Syndrome (Tanzi and Hyman (1991) *Nature* 350:564 and Rumble *et al.* (1989) *N. Engl. J. Med.* 320:1446). *In situ* hybridization studies have shown that APP 717 mutations do not alter APP expression (Harrison *et al.* (1991) *Neurorep.* 2:152).

The V717I (APP 717 Val->Ile), V717G (APP 717 Val->Gly) and V717F (APP 717 Val->Phe) mutations would destabilise a putative stem loop structure and destroy a possible iron-responsive element between base pairs 2131 and 2156 (Tanzi and Hyman, *loc. cit.*). There are several other possible mutations which could also disrupt this structure, many of which would be silent at the protein level; yet these mutations specifically referred to have involved a change to the same amino acid, and no silent changes or changes to other amino acids have been reported prior to the work described herein. Examination of sequence data from 10 other mammalian species (Johnstone *et al.* (1991) *Mol. Brain Res.* 10:299) shows that while the valine residue at codon 717 is conserved in all of them, the putative stem loop structure postulated from the human sequence (Tanzi and Hyman *loc. cit.*) would not be predicted to occur in either cattle or sheep; and in pig and mouse the consensus sequence for the iron-responsive elements is not present. Finally, such stem loop structures are believed to modulate gene translation by altering mRNA stability (Klausner and Harford (1989) *Science* 246:870); however, Harrison and colleagues (Harrison *et al. loc. cit.*) have shown by *in situ* hybridization that APP mRNAs are not grossly altered in the brain of an individual with the V717I mutation. For these reasons, it is believed likely that alterations in the rate of APP translation caused by the

specific mutations identified are not likely to be the key to their pathogenicity.

The fact that the specific mutations discussed involve different changes (Val->Ile, Val->Gly, and Val->Phe) suggests that neither side-chain hydrophobicity nor side-chain bulk is the crucial issue. All examples of APP alleles that encode an amino acid other than valine at codon 717, cosegregate with FAD; suggesting that the valine that occurs at position 717 in wild-type APP770 or APP751 is a critical amino acid residue for non-pathogenic APP proteolytic processing (i.e., by the constitutive cleavage pathway).

The major metabolic pathway for the APP molecule involves cleavage within the beta-amyloid fragment (Esch *et al. loc. cit.*). To generate beta-amyloid, there must be a second pathway in which APP is cleaved outside this sequence. Such a cleavage would be likely to leave a stub of the APP molecule containing the beta-amyloid fragment embedded in the membrane. Possibly, the beta-amyloid-containing fragment which is generated by the second pathway is degraded by peptidase action; the reported mutations may be pathogenic because peptides which contain them may be more resistant to the actions of this peptidase. Therefore, genetic alterations in the APP gene which result in altered (generally reduced) degradative properties are particularly important in the application of the invention. There are several methodologies available from recombinant DNA technology which may be used for detecting and identifying a genetic mutation responsible for Alzheimer's disease. These include direct probing, polymerase chain reaction (PCR) methodology, restriction fragment length polymorphism (RFLP) analysis and single strand conformational analysis (SSCA).

Detection of point mutations using direct probing involves the use of oligonucleotide probes which may be prepared synthetically or by nick translation. The DNA probes may be suitably labelled using, for example, a radiolabel, enzyme label, fluorescent label, biotin-avidin label and the like for subsequent visualization in for example a Southern blot hybridization procedure. The labelled probe is reacted with the sample DNA bound to a nitrocellulose or Nylon 66

substrate. The areas that carry DNA sequences complementary to the labelled DNA probe become labelled themselves as a consequence of the reannealing reaction. The areas of the filter that exhibit such labelling may then be visualized, for example, by autoradiography.

5                   Alternative probing techniques, such as ligase chain reaction (LCR) involve the use of mismatch probes, *i.e.*, probes which have full complementarity with the target except at the point of the mutation. The target sequence is then allowed to hybridize both with oligonucleotides having full complementarity and  
10                   oligonucleotides containing a mismatch, under conditions which will distinguish between the two. By manipulating the reaction conditions it is possible to obtain hybridization only where there is full complementarity. If a mismatch is present then there is significantly reduced hybridization.

                  The polymerase chain reaction (PCR) is a technique that amplifies specific DNA sequences with remarkable efficiency. Repeated cycles of  
15                   denaturation, primer annealing and extension carried out with a heat stable enzyme Taq polymerase leads to exponential increases in the concentration of desired DNA sequences.

                  Given a knowledge of the nucleotide sequence encoding the precursor of amyloid protein of AD (Kang *et al. loc. cit.*, and Yoshikai, above)  
20                   it may be possible to prepare synthetic oligonucleotides complementary to sequences which flank the DNA of interest. Each oligonucleotide is complementary to one of the two strands. The DNA is then denatured at high temperatures (e.g., 95°C) and then reannealed in the presence of a large molar excess of oligonucleotides. The oligonucleotides, oriented with their 3' ends  
25                   pointing towards each other, hybridize to opposite strands of the target sequence and prime enzymatic extension along the nucleic acid template in the presence of the four deoxyribonucleotide triphosphates. The end product is then denatured again for another cycle. After this three-step cycle has been repeated several times, amplification of a DNA segment by more than one million fold can be achieved.

The resulting DNA may then be directly sequenced in order to locate any genetic alteration. Alternatively, it may be possible to prepare oligonucleotides that will only bind to altered DNA, so that PCR will only result in multiplication of the DNA if the mutation is present. Following PCR, allele-specific oligonucleotide hybridization (Dihella et al. (1988) Lancet 1:497) may be used to detect the AD point mutation. Alternatively an adaptation of PCR called amplification of specific alleles (PASA) can be employed; this uses differential amplification for rapid and reliable distinction between alleles that differ at a single base pair.

In yet another method PCR may be followed by restriction endonuclease digestion with subsequent analysis of the resultant products. The substitution of T for C at base pair 2149, found as a result of sequencing exon 17, creates a *Bcl*II restriction site. The creation of this restriction endonuclease recognition site facilitates the detection of the AD mutation using RFLP analysis or by detection of the presence or absence of a polymorphic *Bcl*II site in a PCR product that spans codon 717.

For RFLP analysis, DNA is obtained, for example, from the blood of the subject suspected of having AD and from a normal subject is digested with the restriction endonuclease *Bcl*II and subsequently separated on the basis of size using agarose gel electrophoresis. The Southern blot technique can then be used to detect, by hybridization with labeled probes, the products of endonuclease digestion. The patterns obtained from the Southern blot can then be compared. Using such an approach, DNA spanning an Alzheimer's mutation that creates or removes a restriction site at codon 717, such as the *Bcl*II site, is detected by determining the number of bands detected and comparing this number to a reference allele that has a codon 717 allele that encodes valine.

Correspondingly, the substitution of G for T at base pair 2150 creates a *Sfa*NI restriction site (GCATC), which may be exploited in a manner similar to that described above, *mutatis mutandis*. Similar creation of additional restriction sites by nucleotide substitutions within codon 717, wherein the codon

717 encodes an amino acid other than valine, can be readily calculated by reference to the genetic code and a list of nucleotide sequences recognized by restriction endonucleases (Promega Protocols and Applications Guide, (1991) Promega Corporation, Madison, Wisconsin).

5                   Single strand conformational analysis (SSCA) (Orita *et al.* (1989) *Genomics* 5:874 and Orita *et al.* (1990) *Genomics* 6:271) offers a relatively quick method of detecting sequence changes which may be appropriate in at least some instances.

10                   PCR amplification of specific alleles (PASA) is a rapid method of detecting single-base mutations or polymorphisms (Newton *et al.* (1989) Nucleic Acids Res. 17:2503; Nichols *et al.* (1989) Genomics 5:535; Okayama *et al.* (1989) J. Lab. Clin. Med. 114:105; Sarkar *et al.* (1990) Anal. Biochem. 186:64; Sommer *et al.* (1989) Mayo Clin. Proc. 64:1361; Wu (1989) Proc. Natl. Acad. Sci. U.S.A. 86:2757; and Dutton *et al.* (1991) Biotechniques 11:700). PASA (also known as  
15                   allele specific amplification) involves amplification with two oligonucleotide primers such that one is allele-specific. The desired allele is efficiently amplified, while the other allele(s) is poorly amplified because it mismatches with a base at or near the 3' end of the allele-specific primer. Thus, PASA or the related method of PAMSA may be used to specifically amplify one or more variant APP codon  
20                   717 alleles. Where such amplification is done on genetic material (or RNA) obtained from an individual, it can serve as a method of detecting the presence of one or more variant APP codon 717 alleles in an individual.

25                   Similarly, a method known as a ligase chain reaction (LCR) has been used to successfully detect a single-base substitution in a hemoglobin allele that causes sickle cell anemia (Barany *et al.* (1991) Proc. Natl. Acad. Sci. U.S.A. 88:189; Weiss (1991) Science 254:1292). LCR probes may be combined, or multiplexed for simultaneously screening for multiple different mutations. Thus, one method of screening for variant APP codon 717 alleles is to multiplex at least two, and preferably all, LCR probes that will detect an APP allele having a codon

717 that does not encode valine, but that does encode an amino acid. The universal genetic code provides the degenerate sequences of all the encoded non-valine amino acids, thus LCR probe design for detecting any particular variant codon 717 allele is straightforward, and multiplexed pools of such LCR probes may be selected in the discretion of a practitioner for his particular desired use.

In performing diagnosis using any of the above techniques or variations thereof, it is preferable that several individuals are examined. These may include an unaffected parent, an affected parent, an affected sibling, an unaffected sibling as well as other perhaps more distant family members.

10

#### Model Animals and Cell Lines

Having identified specific mutations in codon 717 of the  $\beta$ -amyloid gene as a cause of familial Alzheimer's disease (FAD), it is possible, using genetic manipulation, to develop transgenic model systems and/or whole cell systems containing the mutated FAD gene (or a portion thereof) for use, for example, as model systems for screening for drugs and evaluating drug effectiveness. Additionally, such model systems provide a tool for defining the underlying biochemistry of APP and  $\beta$ -amyloid metabolism, which thereby provides a basis for rational drug design.

20

One type of cell system can be naturally derived. For this, blood samples from the affected subject must be obtained in order to provide the necessary cells which can be permanently transformed into a lymphoblastoid cell line using, for example, Epstein-Barr virus.

Once established, such cell lines can be grown continuously in suspension culture and may be used for a variety of *in vitro* experiments to study APP expression and processing.

25

Since the FAD mutation is dominant, an alternative method for constructing a cell line is to engineer genetically a mutated gene, or a portion thereof spanning codon 717, into an established (either stably or transiently) cell

line of choice. Sisodia (1990) *Science* 248:492) has described the insertion of a normal APP gene, by transfection, into mammalian cells. Oltersdorf *et al.* ((1990) *J. Biol. Chem.* 265:4492) describe the insertion of APP into immortalized eukaryotic cell lines.

5 Baculovirus expression systems are useful for high level expression of heterologous genes in eukaryotic cells. Knops *et al.* (1991) *J. Biol. Chem.* 266(11):7285 describes the expression of APP using such a system.

10 In yet a further use of the present method, it may be possible to excise the mutated gene (i.e., a variant APP codon 717 gene) for use in the creation of transgenic animals containing the mutated gene. For example, an entire human variant APP codon 717 allele may be cloned and isolated, either in parts or as a whole, in a cloning vector (e.g.,  $\lambda$ Charon35, cosmid, or yeast artificial chromosome). The human variant APP codon 717 gene, either in parts or in whole, may be transferred to a host nonhuman animal, such as a mouse. As  
15 a result of the transfer, the resultant transgenic nonhuman animal will preferably express one or more variant APP codon 717 polypeptides. Most preferably, a transgenic nonhuman animal of the invention will express one or more variant APP codon 717 polypeptides in a neuron-specific manner (Wirak *et al.* (1991) *EMBO* 10:289). This may be accomplished by transferring substantially the entire human  
20 APP gene (encoding a codon 717 mutant) including the 4.5 kilobase sequence that is adjacent to and upstream of the first major APP transcriptional start site.

Alternatively, one may design minigenes encoding variant APP codon 717 polypeptides. Such minigenes may contain a CDNA sequence encoding a variant APP codon 717 polypeptide, preferably full-length, a combination of APP  
25 gene exons, or a combination thereof, linked to a downstream polyadenylation signal sequence and an upstream promoter (and preferably enhancer). Such a minigene construct will, when introduced into an appropriate transgenic host (e.g., mouse or rat), express an encoded variant APP codon 717 polypeptide, most preferably a variant APP codon 717 polypeptide that contains either an isoleucine,



glycine, or phenylalanine residue at codon 717 of APP770 or the corresponding position in an APP isoform or fragment.

One approach to creating transgenic animals is to target a mutation to the desired gene by homologous recombination in an embryonic stem (ES) cell line *in vitro* followed by microinjection of the modified ES cell line into a host blastocyst and subsequent incubation in a foster mother (see Frohman and Martin (1989) Cell 56:145). Alternatively, the technique of microinjection of the mutated gene, or a portion thereof, into a one-cell embryo followed by incubation in a foster mother can be used. Various uses of transgenic animals, particularly transgenic animals that express a wild-type APP isoform or fragment, are disclosed in Wirak *et al.* (1991) EMBO, 10(2):289; Schilling *et al.* (1991) Gene 98(2):225; Quon *et al.* (1991) Nature 352:239; Wirak *et al.* (1991) Science 253:323; and Kawabata *et al.* (1991) Nature 354:476. Additional methods for producing transgenic animals are known in the art.

Alternatively, site-directed mutagenesis and/or gene conversion can be used to mutate a murine (or other nonhuman) APP gene allele, either endogenous or transfected, such that the mutated allele does not encode valine at the codon position in the mouse APP gene that corresponds to codon 717 (of APP770) of the human APP gene (such position is readily identified by homology matching of the murine APP gene or APP protein to the human APP gene or APP770 protein). Preferably, such a mutated murine allele would encode isoleucine or glycine or phenylalanine at the corresponding codon position.

### Therapeutics

Having detected the genetic mutation in the gene sequence coding for  $\beta$ -amyloid protein in an individual not yet showing overt signs of familial AD, using the method of the present invention, it may be possible to employ gene therapy, in the form of gene implants, to prevent the development of the disease.

Additional embodiments directed to modulation of the production of

variant APP proteins include methods that employ specific antisense polynucleotides complementary to all or part of a variant APP sequence, or for some embodiments a wild-type APP sequence. Such complementary antisense polynucleotides may include nucleotide substitutions, additions, deletions, or transpositions, so long as specific hybridization to the relevant target sequence, *i.e.*, a variant APP codon 717 sequence, is retained as a property of the polynucleotide. Thus, an antisense polynucleotide must preferentially bind to a variant APP (*i.e.*, codon 717 does not encode valine) sequence as compared to a wild-type APP (*i.e.*, codon 717 does encode valine). It is evident that the antisense polynucleotide must reflect the exact nucleotide sequence of the variant allele (or wild-type allele where desired) and not a degenerate sequence.

Complementary antisense polynucleotides include soluble antisense RNA or DNA oligonucleotides which can hybridize specifically to a variant APP mRNA species and prevent transcription of the mRNA species and/or translation of the encoded polypeptide (Ching et al. (1989) Proc. Natl. Acad. Sci. U.S.A. 86:10006; Broder et al. (1990) Ann. Int. Med. 113:604; Loreau et al. (1990) FEBS Letters 274:53-56); Holcenberg et al. WO91/11535; U.S. No. 7,530,165 ("New human CRIPTO gene" - publicly available through Derwent Publications Ltd., Rochdale House, 128 Theobalds Road, London, UK); WO91/09865; WO91/04753; WO90/13641; and EP 386563, each of which is incorporated herein by reference). The antisense polynucleotides therefore inhibit production of the variant APP polypeptides. Antisense polynucleotides may preferentially inhibit transcription and/or translation of mRNA corresponding to a variant (or wild-type) polypeptides can inhibit T lymphocyte activation.

Antisense polynucleotides may be produced from a heterologous expression cassette in a transfectant cell or transgenic cell or animal, such as a transgenic neural, glial, or astrocytic cell, preferably where the expression cassette contains a sequence that promotes cell-type specific expression (Wirak et al. *loc. cit.*). Alternatively, the antisense polynucleotides may comprise soluble

oligonucleotides that are administered to the external milieu, either in the culture medium in vitro or in the circulatory system or interstitial fluid in vivo. Soluble antisense polynucleotides present in the external milieu have been shown to gain access to the cytoplasm and inhibit translation of specific mRNA species. In some  
5       embodiments the antisense polynucleotides comprise methylphosphonate moieties. For general methods relating to antisense polynucleotides, see Antisense RNA and DNA, (1988), D.A. Melton, Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY).

10       Mutant APP Antigens and Monoclonal Antibodies

In yet another aspect of the invention, having detected a genetic alteration in a gene sequence coding for APP, it may be possible to obtain samples of the altered  $\beta$ -amyloid protein from the same source. This protein may be derived from the brain tissue of a subject diagnosed as suffering from Alzheimer's  
15       disease, or more preferably are produced by recombinant DNA methods or are synthesized by direct chemical synthesis on a solid support. Such polypeptides will contain an amino acid sequence of an APP variant allele spanning codon 717. Examples of such sequences are:

- (a) -Ile-Ala-Thr-Val-Ile-Gly-Ile-Thr-Leu- [SEQ ID NO:7]
- 20       (b) -Ile-Ala-Thr-Val-Ile-Met-Ile-Thr-Leu- [SEQ ID NO:8]
- (c) -Ile-Ala-Thr-Val-Ile-Ala-Ile-Thr-Leu- [SEQ ID NO:9]
- (d) -Ile-Ala-Thr-Val-Ile-Ser-Ile-Thr-Leu- [SEQ ID NO:10]
- (e) -Ile-Ala-Thr-Val-Ile-Ile-Ile-Thr-Leu- [SEQ ID NO:11]
- (f) -Ile-Ala-Thr-Val-Ile-Leu-Ile-Thr-Leu- [SEQ ID NO:12]
- 25       (g) -Ile-Ala-Thr-Val-Ile-Thr-Ile-Thr-Leu- [SEQ ID NO:13]
- (h) -Ile-Ala-Thr-Val-Ile-Pro-Ile-Thr-Leu- [SEQ ID NO:14]
- (i) -Ile-Ala-Thr-Val-Ile-His-Ile-Thr-Leu- [SEQ ID NO:15]
- (j) -Ile-Ala-Thr-Val-Ile-Cys-Ile-Thr-Leu- [SEQ ID NO:16]
- (k) -Ile-Ala-Thr-Val-Ile-Tyr-Ile-Thr-Leu- [SEQ ID NO:17]

- (l) -Ile-Ala-Thr-Val-Ile-Phe-Ile-Thr-Leu- [SEQ ID NO:18]  
(m) -Ile-Ala-Thr-Val-Ile-Glu-Ile-Thr-Leu- [SEQ ID NO:19]  
(n) -Ile-Ala-Thr-Val-Ile-Trp-Ile-Thr-Leu- [SEQ ID NO:20]  
(o) -Ile-Ala-Thr-Val-Ile-Arg-Ile-Thr-Leu- [SEQ ID NO:21]  
5 (p) -Ile-Ala-Thr-Val-Ile-Asp-Ile-Thr-Leu- [SEQ ID NO:22]  
(q) -Ile-Ala-Thr-Val-Ile-Asn-Ile-Thr-Leu- [SEQ ID NO:23]  
(r) -Ile-Ala-Thr-Val-Ile-Lys-Ile-Thr-Leu- [SEQ ID NO:24]  
(s) -Ile-Ala-Thr-Val-Ile-Gln-Ile-Thr-Leu- [SEQ ID NO:25]

10 Using such polypeptide material it may then be possible to prepare antisera and monoclonal antibodies using, for example, the method of Kohler and Milstein ((1975) *Nature* 256:495). Such monoclonal antibodies could then form the basis of a diagnostic test.

15 Such variant APP polypeptides may be used to immunize an animal for the production of specific antibodies. These antibodies may comprise a polyclonal antiserum or may comprise a monoclonal antibody produced by hybridoma cells. For general methods to prepare antibodies, see Antibodies: A Laboratory Manual, (1988) E. Harlow and D. Lane, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY, which is incorporated herein by reference.

20 For example but not for limitation, a recombinantly produced fragment of a variant APP codon 717 polypeptide can be injected into a mouse along with an adjuvant so as to generate an immune response. Murine immunoglobulins which bind the recombinant fragment with a binding affinity of at least  $1 \times 10^7 \text{ M}^{-1}$  can be harvested from the immunized mouse as an antiserum,  
25 and may be further purified by affinity chromatography or other means. Additionally, spleen cells are harvested from the mouse and fused to myeloma cells to produce a bank of antibody-secreting hybridoma cells. The bank of hybridomas can be screened for clones that secrete immunoglobulins which bind the recombinantly produced fragment with an affinity of at least  $1 \times 10^6 \text{ M}^{-1}$ . More

specifically, immunoglobulins that bind to the variant APP codon 717 polypeptide but have limited crossreactivity with a wild-type (i.e., codon 717 encodes valine) APP polypeptide are selected, either by preabsorption with wild-type APP or by screening of hybridoma cell lines for specific idiotypes that preferentially bind the variant as compared to the wild-type.

The nucleic acid sequences of the present invention capable of ultimately expressing the desired variant APP polypeptides can be formed from a variety of different polynucleotides (genomic or cDNA, RNA, synthetic oligonucleotides, etc.) as well as by a variety of different techniques.

As stated previously, the DNA sequences will be expressed in hosts after the sequences have been operably linked to (i.e., positioned to ensure the functioning of) an expression control sequence. These expression vectors are typically replicable in the host organisms either as episomes or as an integral part of the host chromosomal DNA. Commonly, expression vectors will contain selection markers, e.g., tetracycline resistance or hygromycin resistance, to permit detection and/or selection of those cells transformed with the desired DNA sequences (see, e.g., U.S. Patent 4,704,362, which is incorporated herein by reference).

Polynucleotides encoding a variant APP codon 717 polypeptide may include sequences that facilitate transcription (expression sequences) and translation of the coding sequences, such that the encoded polypeptide product is produced. Construction of such polynucleotides is well known in the art and is described further in Maniatis et al. Molecular Cloning: A Laboratory Manual, 2nd Ed. (1989), Cold Spring Harbor, N.Y. For example, but not for limitation, such polynucleotides can include a promoter, a transcription termination site (polyadenylation site in eukaryotic expression hosts), a ribosome binding site, and, optionally, an enhancer for use in eukaryotic expression hosts, and, optionally, sequences necessary for replication of a vector.

E. coli is one prokaryotic host useful particularly for cloning the DNA sequences of the present invention. Other microbial hosts suitable for use include bacilli, such as Bacillus subtilis, and other enterobacteriaceae, such as Salmonella, Serratia, and various Pseudomonas species. In these prokaryotic hosts, one can also make expression vectors, which will typically contain expression control sequences compatible with the host cell (e.g., an origin of replication). In addition, any number of a variety of well-known promoters will be present, such as the lactose promoter system, a tryptophan (trp) promoter system, a beta-lactamase promoter system, or a promoter system from phage lambda. The promoters will typically control expression, optionally with an operator sequence, and have ribosome binding site sequences and the like, for initiating and completing transcription and translation.

Other microbes, such as yeast, may also be used for expression. Saccharomyces is a preferred host, with suitable vectors having expression control sequences, such as promoters, including 3-phosphoglycerate kinase or other glycolytic enzymes, and an origin of replication, termination sequences and the like as desired.

In addition to microorganisms, mammalian tissue cell culture may also be used to express and produce the polypeptides of the present invention (see, Winnacker, "From Genes to Clones," VCH Publishers, N.Y., N.Y. (1987), which is incorporated herein by reference). Eukaryotic cells are actually preferred, because a number of suitable host cell lines capable of secreting intact human proteins have been developed in the art, and include the CHO cell lines, various COS cell lines, HeLa cells, myeloma cell lines, Jurkat cells, etc. Expression vectors for these cells can include expression control sequences, such as an origin of replication, a promoter, an enhancer (Queen et al. (1986) Immunol. Rev. 89:49, which is incorporated herein by reference), and necessary processing information sites, such as ribosome binding sites, RNA splice sites, polyadenylation sites, and transcriptional terminator sequences. Preferred expression control sequences are

promoters derived from immunoglobulin genes, SV40, Adenovirus, Bovine Papilloma Virus, and the like. The vectors containing the DNA segments of interest (e.g., polypeptides encoding a variant APP polypeptide) can be transferred into the host cell by well-known methods, which vary depending on the type of cellular host. For example, calcium chloride transfection is commonly utilized for prokaryotic cells, whereas calcium phosphate treatment or electroporation may be used for other cellular hosts. (See, generally, Maniatis, et al. Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, (1982), which is incorporated herein by reference.)

Alternatively, homologous recombination may be used to insert an APP mutant sequence into a host genome at a specific site, for example, at a host APP locus. In one type of homologous recombination, one or more host sequence(s) are replaced; for example, a host APP allele (or portion thereof) is replaced with a mutant APP allele (or portion thereof). In addition to such gene replacement methods, homologous recombination may be used to target a mutant APP allele to a specific site other than a host APP locus. Homologous recombination may be used to produce transgenic non-human animals and/or cells that incorporate mutant APP alleles.

The method lends itself readily to the formulation of test kits which can be utilized in diagnosis. Such a kit would comprise a carrier being compartmentalized to receive in close confinement one or more containers wherein a first container may contain suitably labelled DNA probes. Other containers may contain reagents useful in the localization of the labelled probes, such as enzyme substrates. Still other containers may contain a restriction enzyme (such as *BclI*), buffers and the like, together with instructions for use.

#### EXPERIMENTAL EXAMPLES

The following examples are provided for illustration and are not intended to limit the invention to the specific example provided.

EXAMPLE 1 - Detection of a Val->Ile mutation in the  $\beta$ -amyloid (APP) gene

The segregation of AD and markers along the long arm of chromosome 21 in a single family with autopsy-confirmed Alzheimer's disease (see Fig. 1) were examined. DNA samples were available from a total of six affected and 33 unaffected and at risk individuals.

The APP gene in an affected family member was analyzed by polymerase chain reaction (PCR) direct sequencing using intronic primers (Gyllenstein, U. in PCR Technology, Ed. Erlich, H.A., Stockton Press, 45-60, 1989; Yoshikai *et al.* (1990) *Gene* 87:257). (Fig. 2). The primers were made according to the manufacturer's protocol using a Gene Assembler Plus (Pharmacia LKB).

PCR was carried out using the following intronic primers in order to amplify exon 17 of the APP gene:

[A] 5'-CCTCATCCAAATGTCCCCGTCATT-3' [SEQ ID NO:26] AND  
[B] 5'-GCCTAATTCTCTCATAGTCTTAATTCCCAC-3' [SEQ ID NO 27]

PCR conditions were 94°C for 10min to denature; then 35 cycles of 60°C for 1min, 72°C for 3min, 94°C for 1.5min; and a single cycle of 72°C for 10min. The reaction was carried out using 10mM tris-HCl pH 8.3, 50mM potassium chloride, 0.01% gelatin, 1.5mM magnesium chloride, 200 $\mu$ M of dNTPs, 50 pmoles of each PCR primer and 1 unit of Taq polymerase. The total final reaction volume was 25 $\mu$ l.

A second PCR reaction was then performed with a final concentration of 50 pmol of primer [A] and 0.5 pmol of primer [B]. The PCR product was purified on a centricon 100 microconcentrator (Amicon) and used directly for sequencing with the SEQUENASE kit (version 2.0, United States Biochemical Corp.; the word SEQUENASE is a trade mark) following the manufacturer's protocol.



Exon 17 was sequenced first because it encodes part of the  $\beta$ -amyloid peptide and is the site of the mutation (at APP693) leading to Hereditary Cerebral Haemorrhage with Amyloidosis-Dutch Type (HCHWA-D).

5 Sequencing of exon 17 revealed a C to T transition at base pair 2149, causing a valine to isoleucine change at amino acid 717 (Fig. 2 and Fig. 3).

This C to T transition creates a *Bcl*I restriction site enabling detection within the PCR product (Fig. 4). *Bcl*I digests were carried out at 50°C for 2-4 hours, as recommended by the manufacturer, then electrophoresed in 3% agarose.

10 Screening by PCR of 100 unrelated, normal individuals and 14 cases (9 families) of familial late onset disease failed to demonstrate this substitution. Screening of 11 (9 families) cases of early onset familial disease revealed the *Bcl*I restriction site in two affected individuals from an unrelated family. The genetic data show that the disease loci are linked to the missense mutation. Also, failure  
15 to detect this polymorphism in 200 normal chromosomes supports the contention that it is a pathogenic mutation.

The valine to isoleucine substitution occurs within the transmembrane domain two residues from the C-terminus of the  $\beta$ -amyloid peptide. Computer analysis predicts that the substitution makes the transmembrane more  
20 hydrophobic and might thus anchor the protein more firmly within the membrane. The position of the substitution, two residues from the C-terminus of the  $\beta$ -amyloid peptide may be of significance to the origin of the deposited peptide. This finding links Alzheimer's disease to HCHWA-D, a disease in which amyloid deposition is due to a mutation closer to the N-terminus but within the  $\beta$ -amyloid peptide  
25 (Levy *et al. loc. cit.*).

EXAMPLE 2 - Preparation of a cell line containing a defective  $\beta$ -amyloid (APP) gene

10ml of fresh blood are collected from each individual suffering from familial Alzheimer's disease. Lymphocytes are purified from the blood on a Percoll gradient and mixed with Epstein-Barr virus (EBV). The cells are then plated out in medium supplemented with 10% foetal calf serum, antibiotics, glutamine and Cyclosporin A to kill the T lymphocytes. B lymphocytes which are infected by EBV become immortalized and establish a permanent cell line derived from the B cells of the patient.

A lymphoblastoid cell line, AC21, has been deposited with the European Collection of Animal Cell Cultures, Porton Down.

EXAMPLE 3 - Detection of a Val->Gly mutation in the  $\beta$ -amyloid (APP) gene

A pedigree, designated F19 and shown in Fig. 5, which has autopsy-confirmed AD with an onset age of  $59 \pm 4$  years was identified by observing that an allele of the highly polymorphic dinucleotide repeat marker GT12 (D21S210), which is located close to the APP gene, co-segregated with the disease. Linkage analysis gave a peak lod score between the marker and the disease of 3.02 at a recombination fraction of zero, as the following table shows:

Theta	0	0.01	0.05	0.1	0.2	0.3	0.4
Lod	3.02	2.97	2.75	2.47	1.86	1.22	0.6

Lod scores were calculated with seven liability classes modelling age-dependent penetrances from 0.01 to 0.95 with a phenocopy rate of 0.001 and a gene frequency of 0.001 using MLINK from the LINKAGE package (Lathrop *et al.* (1984) *Proc. Natl. Acad. Sci. USA* 81:3443).

APP exon 17 sequences in an affected and an unaffected member of F19 were determined. In the affected member, there was a G->T transition at position 2150, as can be seen from Fig. 6.

The amplification of exon 17 was performed as described in Example 1 above and Chartier-Harlin *et al.* (1991) *Neurosci. Letts.* 129:134, with the following modifications: (a) the amplification primer sequences were:

5                   ATA-ACC-TCA-TCC-AAA-TGT-CCC-C [SEQ ID NO:28] and  
                  GTA-ACC-CAA-GCA-TCA-TGG-AAG-C [SEQ ID NO:29]; and  
(b) the PCR conditions were 94°C/10 minutes then 35 cycles of 60°C/1 minute, 72°C/1 minute, 94°C/1 minute, followed by 72°C/5 minutes.

10                   50 pmol of the second primer were used to generate single stranded product, which was then purified (Chartier-Harlin *et al. loc. cit.*). The purified product was sequenced with the SEQUENASE kit (2.0) (Trade mark; USB) using a primer of sequence:

                  AAA-TGA-AAT-TCT-TCT-AAT-TGC-G [SEQ ID NO:30].

15                   The presence of the T->C transition creates gel artefacts which were resolved by the inclusion of inosine (SEQUENASE kit) in the sequencing reaction.

20                   Direct sequencing of exons 7 and 16 from affected individuals from F19 (Chartier-Harlin *et al. loc. cit.*) shows that these were of normal sequence and SSCA (Orita *et al. loc. cit.*) and Orita *et al.*) failed to identify changes in exons 2, 3, 7, 9, 12, 13 or 15. SSCA of exon 17 detects both APP693 (Levy, *et al. loc. cit.* and Hardy *et al.* (1991) *Lancet* 337:1342-1343) and APP717 Val->Ile under standard screening conditions and, when modified APP717 Val->Gly.

#### EXAMPLE 4 - Production of Transgenic Animals with Mutant APP Allele

25                   Generation of the constructs: The vector plink was constructed by cloning polylinker between the PvuII and EcoRI sites of pBR322 such that the HindIII end of the polylinker was adjacent to the PvuII site. The ligation destroyed both the EcoRI and PvuII sites associated with the pBR322 segments. The 700bp HpaI to EcoRI fragment of pSV2neo (Southern and Berg (1982) J.

5 Mol. Appl. Genet. 1:327) that contains the SV40 polyadenylation signal was cloned into the HpaI to EcoRI sites of pLink to generate pNotSV. The 200 bp XhoI to PstI fragment of pL2 containing the SV40 16S/1gS splice site (Okayama and Berg (1983) Mol. Cell Biol. 3:280) was isolated, blunted with Klenow, then  
10 cloned into the HpaI site of pNotSV to generate pSplice. The 2.3kb NruI to SpeI fragment of pAPP695 containing the coding region of the cDNA for APP (Tanzi *et al.* (1987) *Science* 235:880) was cloned into the NruI to SpeI site of pSplice to generate pd695. The same strategy was used to generate pd751 using the cDNA for APP751 (Tanzi *et al.* (1988) *Nature* 331:528). A variety of promoters have  
15 been inserted into the pd695 and pd751 vectors by using the unique NruI or the HindIII and NruI sites.

Generation of pshAPP695 & pshAPP751: The construct pAmyproBam was generated by cloning the 1.5kb BamHI fragment of the APP  
20 cDNA into the BamHI site of puc19 xHamY. The 700 bp HindIII to Asp718 fragment of the pAmyproBam (similar to the 700 bp BamHI to Asp718 fragment described in Salbaum *et al.* (1988) *EMBO* 7:2807) was cloned into the HindIII to Asp718 sites of pd695 and pd751 to yield pshAPP695 and pshAPP751.

25 pAPP695 and pAPP751: The pAPP695 and pAPP751 vectors were generated by a three-way ligation of the 3.0 kb EcoRI to XhoI fragment of pAmyProBam, the 1.5kb XhoI to SpeI fragment of APP751cDNA, and the SpeI to EcoRI site of pd751.

30 Generation of pNSE751(+47): The pNSE751 (+47) was constructed using a three-way ligation of the HindIII to KpnI fragment of pNSE6 (Forss-Petter *et al.* (1990) *Neuron* 5:187). The KpnI to BstY1 fragment of pNSE6 and a partial BamHI (-47nt relative to the ATG) to HindIII fragment of pAPP751. This resulted in the generation of a KpnI fragment that was cloned into the KpnI

sites of pNSE751(+47). The BstY1/Bam fusion results in the loss of both sites.

Generation of pNSE751: This vector was generated using a four primer two-step PCR protocol (Higuchi *et al.* (1988) *Nucl. Acids Res.* 16:7351) that resulted in a direct fusion of the NSE initiation codon to the APP coding region. Oligonucleotides C2, 1072, 1073, and A2 (see Nucleotide Sequences, *infra.*) were used to generate a PCR product. The KpnI fragment was generated by digestion with the restriction enzyme. The KpnI fragment was used to replace a similar fragment in pNSE751(+47).

Generation of pNSE751-Hardy and pNSE751-Dutch: The Hardy (APP642 Val->Ile of APP695) and Dutch (APP618 Gln->Glu of APP695) mutations were introduced using a four primer two-PCR protocol. Both sets of reactions used the same "outside primers" with the "inside primers" containing the appropriate mutations. This resulted in the generation of BglII to SpeI fragment after digestion, that contained either the Dutch or the Hardy mutation. The BglII to SpeI fragment of pNSE751 was replaced by the mutated fragment to generate the appropriate vector. The presence of the mutation was conformed by sequence analysis of the vectors.

Generation of pNSE751-Hardy and pNSE751-Dutch: The Hardy VI (APP642 V to I), Hardy VG (APP642 V to G), and Dutch (APP618 E to Q) mutations were introduced using the four primer two-step PCR protocol (Higuchi *et al.* (1988)). The Hardy VI mutant was generated using primers 117/738, 922, 923, and 785; Hardy VG mutant was generated using primers 117/738, 1105, 1106, and 785; Dutch mutant was generated using primers 117/738, 1010, 1011, and 785. In all these mutations the 700 bp BglII to SpeI fragment was isolated by digestion of the PCR product with the restriction enzymes, then cloned into the same sites of pNSE751. The mutations were confirmed by sequence analysis.

Generation of pNFH751: The human NFH gene (Lees et al. (1988) *EMBO* 7(7):1947) was isolated from a genomic library using a rat NFH cDNA as a probe (Lieberburg *et al.* (1989) *Proc. Natl. Acids. Res. USA* 86:2463). An SstI fragment was subcloned into the pSK vector. A pair of PCR primers was generated to place a NruI site at the 3' end of the 150 bp amplified fragment immediately upstream of the initiation codon of the NFH gene. The 5' end contains a KpnI site 150nt upstream of the initiation codon. The final construction of pNFH751 was generated by a three-way ligation of the 5.5b HindIII to KpnI fragment of pNFH8.8, the KpnI to NruI PCR generated fragment, and the HindIII to NruI fragment of pd751. The sequence surrounding the PCR generated fusion at the initiation codon was confirmed by sequence analysis. The Dutch and Hardy variants of pNFH751 were generated by substitution of the 600 bp BgIII to SpeI fragment from a sequence confirmed mutated vector for the same fragment of pNFH751. The presence of the mutation was confirmed by hybridization with the mutated oligomer or by sequence analysis.

Generation of pThy751: The pThy751 vector was generated by a three-way ligation. The HindIII to BamHI fragment of pThy8.2 which was isolated from a human genomic library (Chang *et al.* (1985) *Proc. Natl. Acad. Sci. USA* 82:3819), the synthetic fragment ThyAPP, and the HindIII to NruI fragment of pd751.

ThyAPP:

CAGACTGAGATCCCAGAACCCTAGGTCTGACTCTAGGGTCTTGG[SEQ ID NO:31]

Generation of pThyC100: This pThyC100 construct was generated by a three-way ligation. The 3.6kb HindIII to BamHI fragment of pThy8.2, the synthetic fragment ThyAPP2, and the HindIII to BglII fragment of pd751 or pNSE751 Dutch or pNSE751 Hardy were ligated to yield pThyC100.

ThyAPP2:

CAGACTGAGATCCCAGAACCGATCCTAGGTCTGACTCTAGGGTCTTGG  
[SEQ ID NO:32]

5                   The region around the initiation codon was confirmed by sequence analysis.

10                   Preparation of DNA for injection: The transgene for injection was isolated from the corresponding vector of interest for digestion with NotI and gel electrophoresis. The transgene was purified by using the Gene Clean kit (Bio101), then further purified on an Elutip or HPLC purified on a Nucleogen 4000 column.

15                   Microinjection: The transgene was injected at 2-20 mcg/ml into the most convenient pronucleus (usually the male pronucleus) of FVB or B6D2F2 one-cell embryos (Manipulating the Mouse Embryo, B. Hogan, F. Constantini, E. Lacy, Cold Spring Harbor, 1986). The injected embryos were cultured overnight. Embryos that split to the two-cell stage were implanted into pseudo-pregnant female CD1 mice. The mice were weaned at approximately 21 days. Samples of DNA obtained from tail biopsy were analyzed by Southern blot using a transgene specific probe (usually the SV40 3's splice and polyadenylation signal sequences).  
20                   Transgenic mice harboring at least one copy of the transgene were identified.

                  Use of Transgenic Mice: A mouse that expresses the hAPP gene or its variants can be used to test the pathogenesis of amyloid deposition and therapeutic intervention designed to modulate amyloid deposition.

25                   Biochemical analysis of the transgenic mice reveals possible intermediates in the catabolism of APP that are likely precursors to beta-amyloid. This analysis can be carried out in the animal or in primary tissue culture of the expressing cells.

                  The animal can be used to test potential therapeutic agents. The test group of mice is treated with the test compound administered in an appropriate

5 fashion for a set period. At the conclusion of the test period, the animals are assessed behaviourally, biochemically, and histologically for any possible effects of the test compound. The exact protocol depends on the anticipated mechanism of action of the test compound. Compounds that may have utility in treating AD can be identified using this approach.



## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

## (i) APPLICANT:

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(B) STREET: Four Hedges, Castlehill, LLanblethian,

(C) CITY: Cowbridge

(D) STATE: South Glamorgan

(E) COUNTRY: GB

(ii) TITLE OF INVENTION: Test and Model for Alzheimer's Disease

(iii) NUMBER OF SEQUENCES: 44

(iv) COMPUTER READABLE FORM:  
Not Applicable

(v) CURRENT APPLICATION DATA:

APPLICATION NUMBER: WO PCT/GB92/\_\_\_\_\_

## (2) INFORMATION FOR SEQ ID NO:1:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 695 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

Met	Leu	Pro	Gly	Leu	Ala	Leu	Leu	Leu	Leu	Ala	Ala	Trp	Thr	Ala	Arg	1	5	10	15
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Gln	Ile	Ala	Met	Phe	Cys	Gly	Arg	Leu	Asn	Met	His	Met	Asn	Val	Gln	35	40	45	
Asn	Gly	Lys	Trp	Asp	Ser	Asp	Pro	Ser	Gly	Thr	Lys	Thr	Cys	Ile	Asp	50	55	60	
Thr	Lys	Glu	Gly	Ile	Leu	Gln	Tyr	Cys	Gln	Glu	Val	Tyr	Pro	Glu	Leu	65	70	75	80
Gln	Ile	Thr	Asn	Val	Val	Glu	Ala	Asn	Gln	Pro	Val	Thr	Ile	Gln	Asn	85	90	95	
Trp	Cys	Lys	Arg	Gly	Arg	Lys	Gln	Cys	Lys	Thr	His	Pro	His	Phe	Val	100	105	110	
Ile	Pro	Tyr	Arg	Cys	Leu	Val	Gly	Glu	Phe	Val	Ser	Asp	Ala	Leu	Leu	115	120	125	
Val	Pro	Asp	Lys	Cys	Lys	Phe	Leu	His	Gln	Glu	Arg	Met	Asp	Val	Cys	130	135	140	
Glu	Thr	His	Leu	His	Trp	His	Thr	Val	Ala	Lys	Glu	Thr	Cys	Ser	Glu	145	150	155	160
Lys	Ser	Thr	Asn	Leu	His	Asp	Tyr	Gly	Met	Leu	Leu	Pro	Cys	Gly	Ile	165	170	175	
Asp	Lys	Phe	Arg	Gly	Val	Glu	Phe	Val	Cys	Cys	Pro	Leu	Ala	Glu	Glu	180	185	190	
Ser	Asp	Asn	Val	Asp	Ser	Ala	Asp	Ala	Glu	Glu	Asp	Asp	Ser	Asp	Val	195	200	205	
Trp	Trp	Gly	Gly	Ala	Asp	Thr	Asp	Tyr	Ala	Asp	Gly	Ser	Glu	Asp	Lys	210	215	220	

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Val	Val	Glu	Val	Ala	Glu	Glu	Glu	Glu	Val	Ala	Glu	Val	Glu	Glu	Glu	
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Glu	Ala	Asp	Asp	Asp	Glu	Asp	Asp	Glu	Asp	Gly	Asp	Glu	Val	Glu	Glu	
				245					250						255	
Glu	Ala	Glu	Glu	Pro	Tyr	Glu	Glu	Ala	Thr	Glu	Arg	Thr	Thr	Ser	Ile	
			260					265						270		
Ala	Thr	Thr	Thr	Thr	Thr	Thr	Thr	Glu	Ser	Val	Glu	Glu	Val	Val	Arg	
			275				280					285				
Val	Pro	Thr	Thr	Ala	Ala	Ser	Thr	Pro	Asp	Ala	Val	Asp	Lys	Tyr	Leu	
	290					295					300					
Glu	Thr	Pro	Gly	Asp	Glu	Asn	Glu	His	Ala	His	Phe	Gln	Lys	Ala	Lys	
305					310					315					320	
Glu	Arg	Leu	Glu	Ala	Lys	His	Arg	Glu	Arg	Met	Ser	Gln	Val	Met	Arg	
				325					330					335		
Glu	Trp	Glu	Glu	Ala	Glu	Arg	Gln	Ala	Lys	Asn	Leu	Pro	Lys	Ala	Asp	
			340					345					350			
Lys	Lys	Ala	Val	Ile	Gln	His	Phe	Gln	Glu	Lys	Val	Glu	Ser	Leu	Glu	
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	370					375					380					
Arg	Val	Glu	Ala	Met	Leu	Asn	Asp	Arg	Arg	Arg	Leu	Ala	Leu	Glu	Asn	
385					390					395					400	
Tyr	Ile	Thr	Ala	Leu	Gln	Ala	Val	Pro	Pro	Arg	Pro	Arg	His	Val	Phe	
				405					410					415		
Asn	Met	Leu	Lys	Lys	Tyr	Val	Arg	Ala	Glu	Gln	Lys	Asp	Arg	Gln	His	
			420					425					430			
Thr	Leu	Lys	His	Phe	Glu	His	Val	Arg	Met	Val	Asp	Pro	Lys	Lys	Ala	
		435					440					445				
Ala	Gln	Ile	Arg	Ser	Gln	Val	Met	Thr	His	Leu	Arg	Val	Ile	Tyr	Glu	
	450					455					460					
Arg	Met	Asn	Gln	Ser	Leu	Ser	Leu	Leu	Tyr	Asn	Val	Pro	Ala	Val	Ala	
465					470					475					480	
Glu	Glu	Ile	Gln	Asp	Glu	Val	Asp	Glu	Leu	Leu	Gln	Lys	Glu	Gln	Asn	
				485					490					495		
Tyr	Ser	Asp	Asp	Val	Leu	Ala	Asn	Met	Ile	Ser	Glu	Pro	Arg	Ile	Ser	
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Tyr	Gly	Asn	Asp	Ala	Leu	Met	Pro	Ser	Leu	Thr	Glu	Thr	Lys	Thr	Thr	
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## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 751 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

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Met Leu Pro Gly Leu Ala Leu Leu Leu Leu Ala Ala Trp Thr Ala Arg
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      20              25              30

Gln Ile Ala Met Phe Cys Gly Arg Leu Asn Met His Met Asn Val Gln
      35              40              45

Asn Gly Lys Trp Asp Ser Asp Pro Ser Gly Thr Lys Thr Cys Ile Asp
      50              55              60

Thr Lys Glu Gly Ile Leu Gln Tyr Cys Gln Glu Val Tyr Pro Glu Leu
      65              70              75              80

Gln Ile Thr Asn Val Val Glu Ala Asn Gln Pro Val Thr Ile Gln Asn
      85              90              95

Trp Cys Lys Arg Gly Arg Lys Gln Cys Lys Thr His Pro His Phe Val
      100             105             110

Ile Pro Tyr Arg Cys Leu Val Gly Glu Phe Val Ser Asp Ala Leu Leu
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Val Pro Asp Lys Cys Lys Phe Leu His Gln Glu Arg Met Asp Val Cys
      130             135             140

Glu Thr His Leu His Trp His Thr Val Ala Lys Glu Thr Cys Ser Glu
      145             150             155             160

Lys Ser Thr Asn Leu His Asp Tyr Gly Met Leu Leu Pro Cys Gly Ile
      165             170             175

Asp Lys Phe Arg Gly Val Glu Phe Val Cys Cys Pro Leu Ala Glu Glu
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Ser Asp Asn Val Asp Ser Ala Asp Ala Glu Glu Asp Asp Ser Asp Val
      195             200             205

Trp Trp Gly Gly Ala Asp Thr Asp Tyr Ala Asp Gly Ser Glu Asp Lys
      210             215             220

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42

Val 225	Val	Glu	Val	Ala	Glu 230	Glu	Glu	Glu	Val	Ala 235	Glu	Val	Glu	Glu	Glu 240
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Glu	Ala	Glu	Glu 260	Pro	Tyr	Glu	Glu	Ala 265	Thr	Glu	Arg	Thr	Thr 270	Ser	Ile
Ala	Thr	Thr 275	Thr	Thr	Thr	Thr	Thr 280	Glu	Ser	Val	Glu	Glu 285	Val	Val	Arg
Glu	Val 290	Cys	Ser	Glu	Gln	Ala 295	Glu	Thr	Gly	Pro	Cys 300	Arg	Ala	Met	Ile
Ser 305	Arg	Trp	Tyr	Phe	Asp 310	Val	Thr	Glu	Gly	Lys 315	Cys	Ala	Pro	Phe	Phe 320
Tyr	Gly	Gly	Cys	Gly 325	Gly	Asn	Arg	Asn	Asn 330	Phe	Asp	Thr	Glu	Glu 335	Tyr
Cys	Met	Ala	Val 340	Cys	Gly	Ser	Ala	Ile 345	Pro	Thr	Thr	Ala	Ala 350	Ser	Thr
Pro	Asp	Ala 355	Val	Asp	Lys	Tyr	Leu 360	Glu	Thr	Pro	Gly	Asp 365	Glu	Asn	Glu
His	Ala 370	His	Phe	Gln	Lys	Ala 375	Lys	Glu	Arg	Leu	Glu 380	Ala	Lys	His	Arg
Glu 385	Arg	Met	Ser	Gln	Val 390	Met	Arg	Glu	Trp	Glu 395	Glu	Ala	Glu	Arg	Gln 400
Ala	Lys	Asn	Leu	Pro 405	Lys	Ala	Asp	Lys	Lys 410	Ala	Val	Ile	Gln	His 415	Phe
Gln	Glu	Lys	Val 420	Glu	Ser	Leu	Glu	Gln 425	Glu	Ala	Ala	Asn	Glu 430	Arg	Gln
Gln	Leu 435	Val	Glu	Thr	His	Met	Ala 440	Arg	Val	Glu	Ala	Met 445	Leu	Asn	Asp
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Pro 465	Pro	Arg	Pro	Arg	His 470	Val	Phe	Asn	Met	Leu 475	Lys	Lys	Tyr	Val	Arg 480
Ala	Glu	Gln	Lys	Asp 485	Arg	Gln	His	Thr	Leu 490	Lys	His	Phe	Glu	His 495	Val
Arg	Met	Val	Asp 500	Pro	Lys	Lys	Ala	Ala 505	Gln	Ile	Arg	Ser	Gln 510	Val	Met
Thr	His	Leu 515	Arg	Val	Ile	Tyr	Glu 520	Arg	Met	Asn	Gln	Ser	Leu 525	Ser	Leu

Leu Tyr Asn Val Pro Ala Val Ala Glu Glu Ile Gln Asp Glu Val Asp  
 530 535 540  
 Glu Leu Leu Gln Lys Glu Gln Asn Tyr Ser Asp Asp Val Leu Ala Asn  
 545 550 555 560  
 Met Ile Ser Glu Pro Arg Ile Ser Tyr Gly Asn Asp Ala Leu Met Pro  
 565 570 575  
 Ser Leu Thr Glu Thr Lys Thr Thr Val Glu Leu Leu Pro Val Asn Gly  
 580 585 590  
 Glu Phe Ser Leu Asp Asp Leu Gln Pro Trp His Ser Phe Gly Ala Asp  
 595 600 605  
 Ser Val Pro Ala Asn Thr Glu Asn Glu Val Glu Pro Val Asp Ala Arg  
 610 615 620  
 Pro Ala Ala Asp Arg Gly Leu Thr Thr Arg Pro Gly Ser Gly Leu Thr  
 625 630 635 640  
 Asn Ile Lys Thr Glu Glu Ile Ser Glu Val Lys Met Asp Ala Glu Phe  
 645 650 655  
 Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys Leu Val Phe Phe  
 660 665 670  
 Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile Gly Leu Met Val  
 675 680 685  
 Gly Gly Val Val Ile Ala Thr Val Ile Val Ile Thr Leu Val Met Leu  
 690 695 700  
 Lys Lys Lys Gln Tyr Thr Ser Ile His His Gly Val Val Glu Val Asp  
 705 710 715 720  
 Ala Ala Val Thr Pro Glu Glu Arg His Leu Ser Lys Met Gln Gln Asn  
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 Gly Tyr Glu Asn Pro Thr Tyr Lys Phe Phe Glu Gln Met Gln Asn  
 740 745 750

## (2) INFORMATION FOR SEQ ID NO:3:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 770 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

Met	Leu	Pro	Gly	Leu	Ala	Leu	Leu	Leu	Leu	Ala	Ala	Trp	Thr	Ala	Arg	1	5	10	15
Ala	Leu	Glu	Val	Pro	Thr	Asp	Gly	Asn	Ala	Gly	Leu	Leu	Ala	Glu	Pro	20	25	30	
Gln	Ile	Ala	Met	Phe	Cys	Gly	Arg	Leu	Asn	Met	His	Met	Asn	Val	Gln	35	40	45	
Asn	Gly	Lys	Trp	Asp	Ser	Asp	Pro	Ser	Gly	Thr	Lys	Thr	Cys	Ile	Asp	50	55	60	
Thr	Lys	Glu	Gly	Ile	Leu	Gln	Tyr	Cys	Gln	Glu	Val	Tyr	Pro	Glu	Leu	65	70	75	80
Gln	Ile	Thr	Asn	Val	Val	Glu	Ala	Asn	Gln	Pro	Val	Thr	Ile	Gln	Asn	85	90	95	
Trp	Cys	Lys	Arg	Gly	Arg	Lys	Gln	Cys	Lys	Thr	His	Pro	His	Phe	Val	100	105	110	
Ile	Pro	Tyr	Arg	Cys	Leu	Val	Gly	Glu	Phe	Val	Ser	Asp	Ala	Leu	Leu	115	120	125	
Val	Pro	Asp	Lys	Cys	Lys	Phe	Leu	His	Gln	Glu	Arg	Met	Asp	Val	Cys	130	135	140	
Glu	Thr	His	Leu	His	Trp	His	Thr	Val	Ala	Lys	Glu	Thr	Cys	Ser	Glu	145	150	155	160
Lys	Ser	Thr	Asn	Leu	His	Asp	Tyr	Gly	Met	Leu	Leu	Pro	Cys	Gly	Ile	165	170	175	
Asp	Lys	Phe	Arg	Gly	Val	Glu	Phe	Val	Cys	Cys	Pro	Leu	Ala	Glu	Glu	180	185	190	
Ser	Asp	Asn	Val	Asp	Ser	Ala	Asp	Ala	Glu	Glu	Asp	Asp	Ser	Asp	Val	195	200	205	
Trp	Trp	Gly	Gly	Ala	Asp	Thr	Asp	Tyr	Ala	Asp	Gly	Ser	Glu	Asp	Lys	210	215	220	



45

Val 225	Val	Glu	Val	Ala	Glu 230	Glu	Glu	Glu	Val	Ala 235	Glu	Val	Glu	Glu	Glu 240
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Ala	Thr	Thr 275	Thr	Thr	Thr	Thr	Thr 280	Glu	Ser	Val	Glu	Glu 285	Val	Val	Arg
Glu	Val 290	Cys	Ser	Glu	Gln	Ala 295	Glu	Thr	Gly	Pro	Cys 300	Arg	Ala	Met	Ile
Ser 305	Arg	Trp	Tyr	Phe	Asp 310	Val	Thr	Glu	Gly	Lys 315	Cys	Ala	Pro	Phe	Phe 320
Tyr	Gly	Gly	Cys	Gly 325	Gly	Asn	Arg	Asn	Asn 330	Phe	Asp	Thr	Glu	Glu 335	Tyr
Cys	Met	Ala	Val	Cys	Gly	Ser	Ala	Met 345	Ser	Gln	Ser	Leu	Leu 350	Lys	Thr
Thr	Gln	Glu	Pro	Leu	Ala	Arg	Asp 360	Pro	Val	Lys	Leu	Pro 365	Thr	Thr	Ala
Ala	Ser 370	Thr	Pro	Asp	Ala	Val 375	Asp	Lys	Tyr	Leu	Glu 380	Thr	Pro	Gly	Asp
Glu 385	Asn	Glu	His	Ala	His 390	Phe	Gln	Lys	Ala	Lys 395	Glu	Arg	Leu	Glu	Ala 400
Lys	His	Arg	Glu	Arg	Met	Ser	Gln	Val	Met 410	Arg	Glu	Trp	Glu	Glu 415	Ala
Glu	Arg	Gln	Ala	Lys	Asn	Leu	Pro	Lys 425	Ala	Asp	Lys	Lys	Ala	Val	Ile
Gln	His	Phe	Gln	Glu	Lys	Val	Glu 440	Ser	Leu	Glu	Gln 445	Glu	Ala	Ala	Asn
Glu 450	Arg	Gln	Gln	Leu	Val	Glu	Thr	His	Met	Ala	Arg 460	Val	Glu	Ala	Met
Leu 465	Asn	Asp	Arg	Arg	Arg 470	Leu	Ala	Leu	Glu	Asn 475	Tyr	Ile	Thr	Ala	Leu 480
Gln	Ala	Val	Pro	Pro	Arg	Pro	Arg	His	Val 490	Phe	Asn	Met	Leu	Lys 495	Lys
Tyr	Val	Arg	Ala	Glu	Gln	Lys	Asp 505	Arg	Gln	His	Thr	Leu	Lys 510	His	Phe
Glu	His	Val	Arg	Met	Val	Asp	Pro 520	Lys	Lys	Ala	Ala	Gln	Ile	Arg	Ser
	515											525			

Gln Val Met Thr His Leu Arg Val Ile Tyr Glu Arg Met Asn Gln Ser  
 530 535 540

Leu Ser Leu Leu Tyr Asn Val Pro Ala Val Ala Glu Glu Ile Gln Asp  
 545 550 555 560

Glu Val Asp Glu Leu Leu Gln Lys Glu Gln Asn Tyr Ser Asp Asp Val  
 565 570 575

Leu Ala Asn Met Ile Ser Glu Pro Arg Ile Ser Tyr Gly Asn Asp Ala  
 580 585 590

Leu Met Pro Ser Leu Thr Glu Thr Lys Thr Thr Val Glu Leu Leu Pro  
 595 600 605

Val Asn Gly Glu Phe Ser Leu Asp Asp Leu Gln Pro Trp His Ser Phe  
 610 615 620

Gly Ala Asp Ser Val Pro Ala Asn Thr Glu Asn Glu Val Glu Pro Val  
 625 630 635 640

Asp Ala Arg Pro Ala Ala Asp Arg Gly Leu Thr Thr Arg Pro Gly Ser  
 645 650 655

Gly Leu Thr Asn Ile Lys Thr Glu Glu Ile Ser Glu Val Lys Met Asp  
 660 665 670

Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys Leu  
 675 680 685

Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile Gly  
 690 695 700

Leu Met Val Gly Gly Val Val Ile Ala Thr Val Ile Val Ile Thr Leu  
 705 710 715 720

Val Met Leu Lys Lys Lys Gln Tyr Thr Ser Ile His His Gly Val Val  
 725 730 735

Glu Val Asp Ala Ala Val Thr Pro Glu Glu Arg His Leu Ser Lys Met  
 740 745 750

Gln Gln Asn Gly Tyr Glu Asn Pro Thr Tyr Lys Phe Phe Glu Gln Met  
 755 760 765

Gln Asn  
 770

## (2) INFORMATION FOR SEQ ID NO:4:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2088 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

ATGCTGCCCCG GTTTGGCACT GCTCCTGCTG GCCGCCTGGA CGGCTCGGGC GCTGGAGGTA	60
CCCCTGATG GTAATGCTGG CCTGCTGGCT GAACCCCAAGA TTGCCATGTT CTGTGGCAGA	120
CTGAACATGC ACATGAATGT CCAGAATGGG AAGTGGGATT CAGATCCATC AGGGACCAAA	180
ACCTGCATTG ATACCAAGGA AGGCATCCTG CAGTATTGCC AAGAAGTCTA CCCTGAACTG	240
CAGATCACCA ATGTGGTAGA AGCCAACCAA CCAGTGACCA TCCAGAACTG GTGCAAGCGG	300
GGCCGCAAGC AGTGCAAGAC CCATCCCCAC TTTGTGATTC CCTACCGCTG CTTAGTTGGT	360
GAGTTTGTA GTGATGCCCT TCTCGTTCCT GACAAGTGCA AATTCTTACA CCAGGAGAGG	420
ATGGATGTTT GCGAAACTCA TCTTCACTGG CACACCGTCG CCAAAGAGAC ATGCAGTGAG	480
AAGAGTACCA ACTTGCATGA CTACGGCATG TTGCTGCCCT GCGGAATTGA CAAGTTCCGA	540
GGGGTAGAGT TTGTGTGTTG CCCACTGGCT GAAGAAAGTG ACAATGTGGA TTCTGCTGAT	600
GCGGAGGAGG ATGACTCGGA TGTCTGGTGG GGCGGAGCAG ACACAGACTA TGCAGATGGG	660
AGTGAAGACA AAGTAGTAGA AGTAGCAGAG GAGGAAGAAG TGGCTGAGGT GGAAGAAGAA	720
GAAGCCGATG ATGACGAGGA CGATGAGGAT GGTGATGAGG TAGAGGAAGA GGCTGAGGAA	780
CCCTACGAAG AAGCCACAGA GAGAACCACC AGCATTGCCA CCACCACCAC CACCACCACA	840
GAGTCTGTGG AAGAGGTGGT TCGAGTTCCT ACAACAGCAG CCAGTACCCC TGATGCCGTT	900
GACAAGTATC TCGAGACACC TGGGGATGAG AATGAACATG CCCATTTCCA GAAAGCCAAA	960
GAGAGGCTTG AGGCCAAGCA CCGAGAGAGA ATGTCCCAGG TCATGAGAGA ATGGGAAGAG	1020
GCAGAACGTC AAGCAAAGAA CTTGCCTAAA GCTGATAAGA AGGCAGTTAT CCAGCATTTT	1080
CAGGAGAAAG TGGAATCTTT GGAACAGGAA GCAGCCAACG AGAGACAGCA GCTGGTGGAG	1140
ACACACATGG CCAGAGTGGA AGCCATGCTC AATGACCGCC GCCGCCTGGC CCTGGAGAAC	1200
TACATCACCG CTCTGCAGGC TGTTCTCCTT CGGCCTCGTC ACGTGTTCAA TATGCTAAAG	1260

AAGTATGTCC	GCGCAGAACA	GAAGGACAGA	CAGCACACCC	TAAAGCATTT	CGAGCATGTG	1320
CGCATGGTGG	ATCCCAAGAA	AGCCGCTCAG	ATCCGGTCCC	AGGTTATGAC	ACACCTCCGT	1380
GTGATTTATG	AGCGCATGAA	TCAGTCTCTC	TCCCTGCTCT	ACAACGTGCC	TGCAGTGGCC	1440
GAGGAGATTC	AGGATGAAGT	TGATGAGCTG	CTTCAGAAAG	AGCAAAACTA	TTCAGATGAC	1500
GTCTTGGCCA	ACATGATTAG	TGAACCAAGG	ATCAGTTACG	GAAACGATGC	TCTCATGCCA	1560
TCTTTGACCG	AAACGAAAAC	CACCGTGGAG	CTCCTTCCCG	TGAATGGAGA	GTTCAGCCTG	1620
GACGATCTCC	AGCCGTGGCA	TTCTTTTGGG	GCTGACTCTG	TGCCAGCCAA	CACAGAAAAC	1680
GAAGTTGAGC	CTGTTGATGC	CCGCCCTGCT	GCCGACCGAG	GACTGACCAC	TCGACCAGGT	1740
TCTGGGTTGA	CAAATATCAA	GACGGAGGAG	ATCTCTGAAG	TGAAGATGGA	TGCAGAATTC	1800
CGACATGACT	CAGGATATGA	AGTTCATCAT	CAAAAATTGG	TGTTCTTTGC	AGAAGATGTG	1860
GGTTCAAACA	AAGGTGCAAT	CATTGGACTC	ATGGTGGGCG	GTGTTGTCAT	AGCGACAGTG	1920
ATCGTCATCA	CCTTGGTGAT	GCTGAAGAAG	AAACAGTACA	CATCCATTCA	TCATGGTGTG	1980
GTGGAGGTTG	ACGCCGCTGT	CACCCCAGAG	GAGCGCCACC	TGTCCAAGAT	GCAGCAGAAC	2040
GGCTACGAAA	ATCCAACCTA	CAAGTTCTTT	GAGCAGATGC	AGAACTAG		2088

## (2) INFORMATION FOR SEQ ID NO:5:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2265 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

ATGCTGCCCCG	GTTTGGCACT	GCTCCTGCTG	GCCGCCTGGA	CGGCTCGGGC	GCTGGAGGTA	60
CCCACTGATG	GTAATGCTGG	CCTGCTGGCT	GAACCCCA	TTGCCATGTT	CTGTGGCAGA	120
CTGAACATGC	ACATGAATGT	CCAGAATGGG	AAGTGGGATT	CAGATCCATC	AGGGACCAAA	180
ACCTGCATTG	ATACCAAGGA	AGGCATCCTG	CAGTATTGCC	AAGAAGTCTA	CCCTGAACTG	240
CAGATCACCA	ATGTGGTAGA	AGCCAACCAA	CCAGTGACCA	TCCAGAAGT	GTGCAAGCGG	300
GGCCGCAAGC	AGTGCAAGAC	CCATCCCCAC	TTTGTGATTC	CCTACCGCTG	CTTAGTTGGT	360
GAGTTTGTA	GTGATGCCCT	TCTCGTTCCT	GACAAGTGCA	AATTCTTACA	CCAGGAGAGG	420
ATGGATGTTT	GCGAAACTCA	TCTTCACTGG	CACACCGTCG	CCAAAGAGAC	ATGCAGTGAG	480
AAGAGTACCA	ACTTGCATGA	CTACGGCATG	TTGCTGCCCT	GCGGAATTGA	CAAGTTCGGA	540
GGGGTAGAGT	TTGTGTGTTG	CCCACTGGCT	GAAGAAAGTG	ACAATGTGGA	TTCTGCTGAT	600
GCGGAGGAGG	ATGACTCGGA	TGTCTGGTGG	GGCGGAGCAG	ACACAGACTA	TGCAGATGGG	660
AGTGAAGACA	AAGTAGTAGA	AGTAGCAGAG	GAGGAAGAAG	TGGCTGAGGT	GGAAGAAGAA	720
GAAGCCGATG	ATGACGAGGA	CGATGAGGAT	GGTGATGAGG	TAGAGGAAGA	GGCTGAGGAA	780
CCCTACGAAG	AAGCCACAGA	GAGAACCACC	AGCATTGCCA	CCACCACCAC	CACCACCACA	840
GAGTCTGTGG	AAGAGGTGGT	TCGAGAGGTG	TGCTCTGAAC	AAGCCGAGAC	GGGGCCGTGC	900
CGAGCAATGA	TCTCCCGCTG	GTACTTTGAT	GTGACTGAAG	GGAAGTGTGC	CCCATTTCTT	960
TACGGCGGAT	GTGGCGGCAA	CCGGAACAAC	CGGAACAAC	TTGACACAGA	AGAGTACTGC	1020
ATGGCCGTGT	GTGGCAGCGC	CATTCCTACA	ACAGCAGCCA	GTACCCCTGA	TGCCGTTGAC	1080
AAGTATCTCG	AGACACCTGG	GGATGAGAAT	GAACATGCCC	ATTTCCAGAA	AGCCAAAGAG	1140
AGGCTTGAGG	CCAAGCACCG	AGAGAGAATG	TCCCAGGTCA	TGAGAGAATG	GGAAGAGGCA	1200
GAACGTCAAG	CAAAGAACTT	GCCTAAAGCT	GATAAGAAGG	CAGTTATCCA	GCATTTCCAG	1260

GAGAAAGTGG	AATCTTTGGA	ACAGGAAGCA	GCCAACGAGA	GACAGCAGCT	GGTGGAGACA	1320
CACATGGCCA	GAGTGGAAGC	CATGCTCAAT	GACCGCCGCC	GCCTGGCCCT	GGAGAACTAC	1380
ATCACCGCTC	TGCAGGCTGT	TCCTCCTCGG	CCTCGTCACG	TGTTCAATAT	GCTAAAGAAG	1440
TATGTCCGCG	CAGAACAGAA	GGACAGACAG	CACACCCTAA	AGCATTTCTGA	GCATGTGCGC	1500
ATGGTGGATC	CCAAGAAAGC	CGCTCAGATC	CGGTCCCAGG	TTATGACACA	CCTCCGTGTG	1560
ATTTATGAGC	GCATGAATCA	GTCTCTCTCC	CTGCTCTACA	ACGTGCCTGC	AGTGGCCGAG	1620
GAGATTCAGG	ATGAAGTTGA	TGAGCTGCTT	CAGAAAGAGC	AAAACCTATTC	AGATGACGTC	1680
TTGGCCAACA	TGATTAGTGA	ACCAAGGATC	AGTTACGGAA	ACGATGCTCT	CATGCCATCT	1740
TTGACCGAAA	CGAAAACCAC	CGTGGAGCTC	CTTCCCCTGA	ATGGAGAGTT	CAGCCTGGAC	1800
GATCTCCAGC	CGTGGCATTTC	TTTTGGGGCT	GACTCTGTGC	CAGCCAACAC	AGAAAACGAA	1860
GTTGAGCCTG	TTGATGCCCCG	CCCTGCTGCC	GACCGAGGAC	TGACCACTCG	ACCAGGTTCT	1920
GGGTTGACAA	ATATCAAGAC	GGAGGAGATC	TCTGAAGTGA	AGATGGATGC	AGAATTCCGA	1980
CATGACTCAG	GATATGAAGT	TCATCATCAA	AAATTGGTGT	TCTTTGCAGA	AGATGTGGGT	2040
TCAAACAAAG	GTGCAATCAT	TGGACTCATG	GTGGGCGGTG	TTGTCATAGC	GACAGTGATC	2100
GTCATCACCT	TGGTGATGCT	GAAGAAGAAA	CAGTACACAT	CCATTCATCA	TGGTGTGGTG	2160
GAGGTTGACG	CCGCTGTCAC	CCCAGAGGAG	CGCCACCTGT	CCAAGATGCA	GCAGAACGGC	2220
TACGAAAATC	CAACCTACAA	GTTCTTTGAG	CAGATGCAGA	ACTAG		2265

## (2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Ile Ala Thr Val Ile Xaa Ile Thr Leu  
1                            5

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 9 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Ile Ala Thr Val Ile Gly Ile Thr Leu  
1 5



53

## (2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Ile Ala Thr Val Ile Met Ile Thr Leu  
1                                    5

(A) LENGTH: 9 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

Ile Ala Thr Val Ile Ala Ile Thr Leu  
1 5

## (2) INFORMATION FOR SEQ ID NO:10:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 9 amino acids

(B) TYPE: amino acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Ile Ala Thr Val Ile Ser Ile Thr Leu  
1 5

## (2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

Ile Ala Thr Val Ile Ile Ile Thr Leu  
1                    5

(2) INFORMATION FOR SEQ ID NO:12:

**(i) SEQUENCE CHARACTERISTICS:**

- (A) LENGTH: 9 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Ile Ala Thr Val Ile Leu Ile Thr Leu  
1 5

(A) LENGTH: 9 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Ile Ala Thr Val Ile Thr Ile Thr Leu  
1 5

Ile Ala Thr Val Ile Pro Ile Thr Leu  
1 5

## (2) INFORMATION FOR SEQ ID NO:15:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

Ile Ala Thr Val Ile His Ile Thr Leu  
1                    5



## (2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

Ile Ala Thr Val Ile Cys Ile Thr Leu  
1                    5

## (2) INFORMATION FOR SEQ ID NO:17:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 9 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

Ile Ala Thr Val Ile Tyr Ile Thr Leu  
1                    5

## (2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

Ile Ala Thr Val Ile Phe Ile Thr Leu  
1                    5

(2) INFORMATION FOR SEQ ID NO:19:

(i) SEQUENCE CHARACTERISTICS:

- SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 9 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

Ile Ala Thr Val Ile Glu Ile Thr Leu  
1 5

65

(2) INFORMATION FOR SEQ ID NO:20:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 9 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

Ile Ala Thr Val Ile Trp Ile Thr Leu  
1 5

66.

## (2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 9 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

Ile Ala Thr Val Ile Arg Ile Thr Leu  
1                    5

67

## (2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

Ile Ala Thr Val Ile Asp Ile Thr Leu  
1                    5

(2) INFORMATION FOR SEQ ID NO:23:

(i) SEQUENCE CHARACTERISTICS:

- SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 9 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

Ile Ala Thr Val Ile Asn Ile Thr Leu  
1 5



69

## (2) INFORMATION FOR SEQ ID NO:24:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 9 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

Ile Ala Thr Val Ile Lys Ile Thr Leu  
1                    5

(A) LENGTH: 9 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

Ile Ala Thr Val Ile Gln Ile Thr Leu  
1 5

71

(2) INFORMATION FOR SEQ ID NO:26:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 24 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (Primer)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

CCTCATCCAA ATGTCCCCGT CATT

24

72

## (2) INFORMATION FOR SEQ ID NO:27:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 30 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: DNA (Primer)

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

GCCTAATTCT CTCATAGTCT TAATTCCCAC

30

73

## (2) INFORMATION FOR SEQ ID NO:28:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 22 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (Primer)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

ATAACCTCAT CCAAATGTCC CC

22

74

## (2) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 22 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (Primer)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

GTAACCCAAG CATCATGGAA GC

22

75

## (2) INFORMATION FOR SEQ ID NO:30:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 22 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (Primer)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

AAATGAAATT CTTCTAATTG CG

22

76

## (2) INFORMATION FOR SEQ ID NO:31:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 44 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

CAGACTGAGA TCCCAGAACC CTAGGTCTGA CTCTAGGGTC TTGG

44



77

## (2) INFORMATION FOR SEQ ID NO:32:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 48 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (Primer)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

CAGACTGAGA TCCAGAACC GATCCTAGGT CTGACTCTAG GGTCTTGG

48

78

## (2) INFORMATION FOR SEQ ID NO:33:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 25 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

CGACCAGGTT GTGGGTGAC AAATA

25

79

## (2) INFORMATION FOR SEQ ID NO:34:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 29 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

AATCTATTCA TGCCTAGTT TGATACAGC

29

80

## (2) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 21 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

ACAGTGATCA TCATCACCTT G

21

81

## (2) INFORMATION FOR SEQ ID NO:36:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 21 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

CAAGGTGATG ATGATCACTG T

21

82

## (2) INFORMATION FOR SEQ ID NO:37:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 28 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

AGCGACAGTG ATCGGCATCA CCTTGGTG

28

83

## (2) INFORMATION FOR SEQ ID NO:38:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 28 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

CACCCAGGTG ATGCCGATCA CTGTCGCT

28

84

## (2) INFORMATION FOR SEQ ID NO:39:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 24 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

ACCCACATCT TGTGCAAAGA ACAC

24



85

## (2) INFORMATION FOR SEQ ID NO:40:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 24 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

GTGTTCTTTG CACAAGATGT GGGT

24

86

## (2) INFORMATION FOR SEQ ID NO:41:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 27 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: DNA (genomic)

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

CCCAGCCATC ATGCTGCCCCG GGTTGGC

27

87

## (2) INFORMATION FOR SEQ ID NO:42:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 32 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:42:

GCCAACCCGG GCAGCATGAT GACTGGGATC TC

32

88

## (2) INFORMATION FOR SEQ ID NO:43:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 21 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:43:

ACCTGCCACT ATACTGGAAT A

21

89

## (2) INFORMATION FOR SEQ ID NO:44:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 21 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:44:

TGTGCATGTT CAGTCTGCCA C

21

WHAT IS CLAIMED IS:

1. An isolated polynucleotide comprising a nucleic acid sequence encoding a codon 717 mutant of human amyloid precursor protein 770 (APP770),  
5 or an isoform or fragment of APP770 having a mutant amino acid residue at the position encoded by codon 717.

2. An isolated polynucleotide of claim 1, wherein the amino acid at the position encoded by codon 717 is an isoleucine, glycine, or phenylalanine.  
10

3. An isolated polynucleotide of claim 1, wherein the nucleic acid sequence is a cDNA.

4. A composition comprising a polynucleotide probe capable of specifically hybridizing to an amyloid precursor protein 770 (APP770) allele exhibiting a mutation at codon 717.  
15

5. A composition of claim 4, wherein codon 717 of the mutant allele encodes an isoleucine, phenylalanine, or glycine.  
20

6. A composition of claim 4, wherein the probe is labeled.

7. A composition of claim 4, wherein the probe comprises at least about 10 nucleotides spanning amino acid 717 of the APP770 allele.  
25

8. A transgenic host comprising a nucleic acid segment encoding a position 717 mutant of human amyloid precursor protein 770 (APP770), or an APP770 isoform or fragment of APP770 having the mutation.

9. A host of claim 8, which is a primary or immortalized eukaryotic cell line.

10. A host of claim 8, which is a bacterium.

5

11. A host of claim 8, wherein the segment is integrated into the host genome.

12. A host of claim 8, which is a non-human animal having the DNA segment incorporated into its germline and which is capable of expressing the mutant APP770 protein.

10

13. A host of claim 12, wherein the mutant APP770 protein is the sole APP770 protein produced by the animal.

15

14. A transgenic non-human animal with germ cells or somatic cells comprising a heterologous gene encoding a position 717 mutant amyloid precursor protein 770 (APP770), which gene upon expression promotes neuropathological characteristics of Alzheimer's disease in the animal.

20

15. A cultured human primary or immortalized cell, comprising a nucleic acid segment encoding a position 717 mutant of human amyloid precursor protein 770 (APP770), or an APP770 isoform or fragment of APP770 having the mutation.

25

16. A method of screening for an agent capable of treating Alzheimer's disease, comprising:

contacting a host of claim 8 with the agent; and monitoring expression or processing of proteins encoded by the mutant APP770 gene.

17. A diagnostic method for determining an inherited predisposition to Alzheimer's disease in a subject, comprising detecting in the subject the presence of an allele of amyloid precursor protein (APP), an isoform or fragment thereof, wherein said allele has a sequence polymorphism at a position  
5 corresponding to codon 717 of APP770.

18. A method of claim 17, wherein said sequence polymorphism is a nucleotide substitution, whereby an isoleucine or glycine is substituted at  
10 codon 717 of APP770.

19. A method of claim 17, wherein said sequence polymorphism is a single nucleotide substitution.

20. A method according to claim 17, wherein the detecting step  
15 comprises sequencing a genomic DNA segment from chromosome 21 of the subject.

21. A method according to claim 17, wherein the detecting step  
20 comprises (i) mixing a nucleic acid sample from the subject with one or more polynucleotide probes capable of hybridizing selectively to an APP gene allele in a reaction and (ii) monitoring the reaction to determine the presence of the gene allele in the sample, thereby indicating whether the subject is at risk for Alzheimer's disease.

22. A method according to claim 20, wherein one probe is a  
25 polynucleotide comprising a sequence of at least about 10 nucleotides spanning codon 717 of an APP770.



23. A method according to claim 22, wherein the probes are oligonucleotides capable of priming polynucleotide synthesis in a polymerase chain reaction, wherein a reaction product comprises a sequence of at least 25 contiguous nucleotides from exon 17 of the APP gene.

5

24. A method according to claim 21, wherein at least one oligonucleotide specifically hybridizes to a sequence present in an intron or flanking region of an APP770 gene.

10

25. A method according to claim 21, wherein the monitoring step comprises analyzing sequencing gel reaction products from the PCR reaction.

15

26. A method according to claim 21, wherein the monitoring step comprises analyzing an autoradiograph of a BclI digest of reaction products from the PCR reaction.

20

27. A method according to claim 17, wherein the detecting step comprises (i) mixing in an immunological assay an APP770 or isoform protein sample from the subject with an antibody reagent specific for the allele and (ii) monitoring the assay to determine specific binding between the antibody reagent and the protein sample, thereby indicating whether the subject is at risk for Alzheimer's disease.

25

28. A method according to claim 27, wherein the antibody reagent is a monoclonal antibody specifically reactive with an antigenic determinant specific for an allele.

30

29. A method for genetic analysis of a human subject which comprises detecting the presence or absence of at least one polymorphism at codon 717 of an APP770 gene of an amyloid precursor protein (APP) gene in the subject.

30. A method according to claim 29, wherein the polymorphism is detected by digesting genomic DNA from the subject with at least one restriction endonuclease and hybridizing resulting fragments with a detecting probe

5 31. A composition comprising a polypeptide free from human proteins, comprising a core sequence:

Ile-Ala-Thr-Val-Ile-X-Ile-Thr-Leu- [SEQ ID NO:6]

wherein X is any of the twenty conventional amino acids except valine.

10 32. A transgenic nonhuman animal containing a polypeptide of claim 31.

33. A transgenic nonhuman animal of Claim 32, wherein the polypeptide is present in the brain.

15 34. An isolated polynucleotide, comprising a nucleic acid sequence encoding a mutant human APP allele that cosegregates with a genetic predisposition to Alzheimer's disease.

20 35. An isolated polypeptide of Claim 34, wherein said mutant human APP allele comprises a codon 717 mutant.

25 36. A method of determining a genetic predisposition of a subject to Alzheimer's disease, the method comprising detecting in the subject's DNA the presence of an allele of a gene encoding amyloid precursor protein (APP).

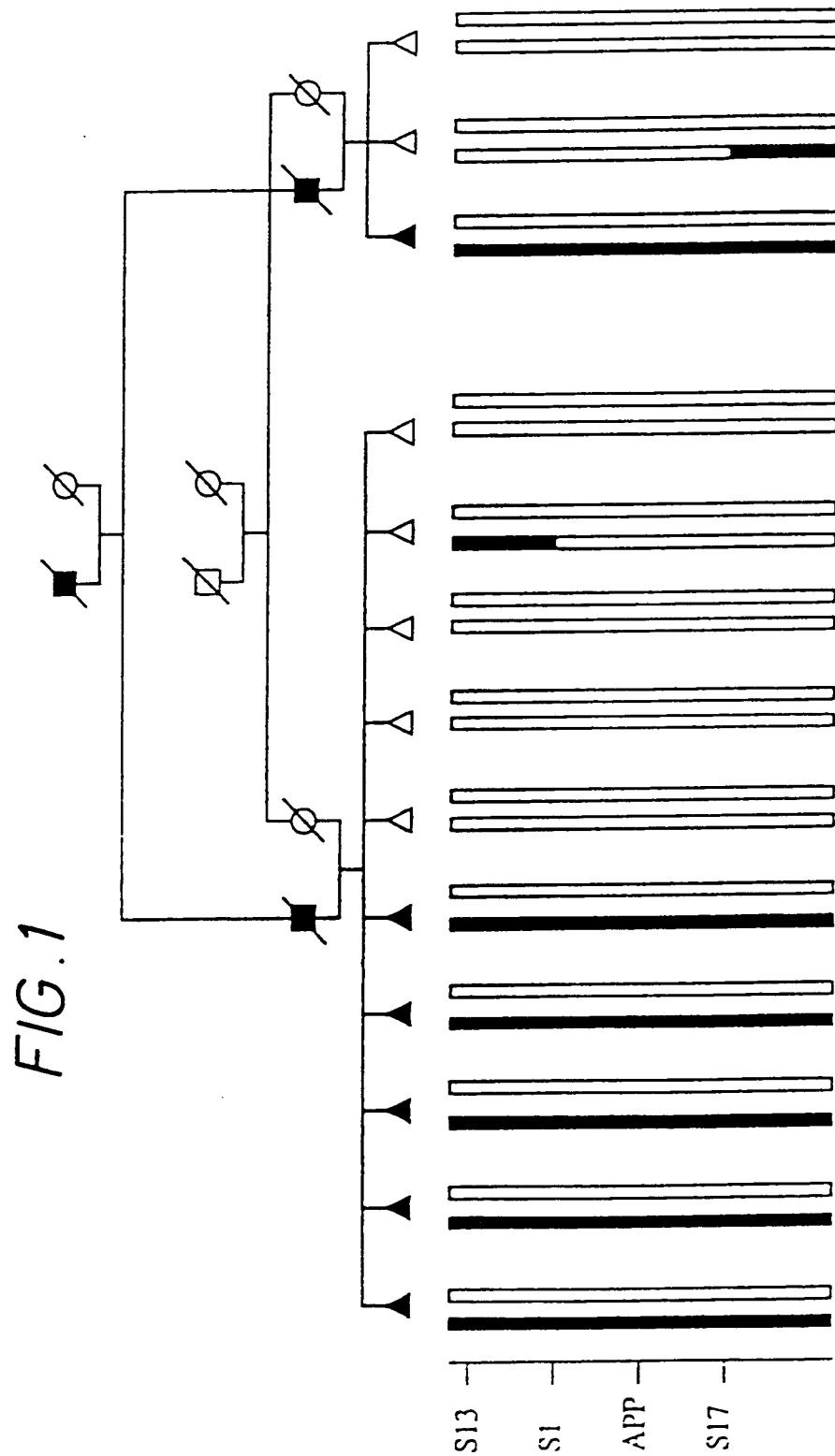
37. A method as claimed in claim 36, wherein the step of detection is carried out on material removed from, and not returned to, the subject's body.

38. A method as claimed in claim 36 or 37, wherein the allele of the gene encodes a substitution mutant of APP.

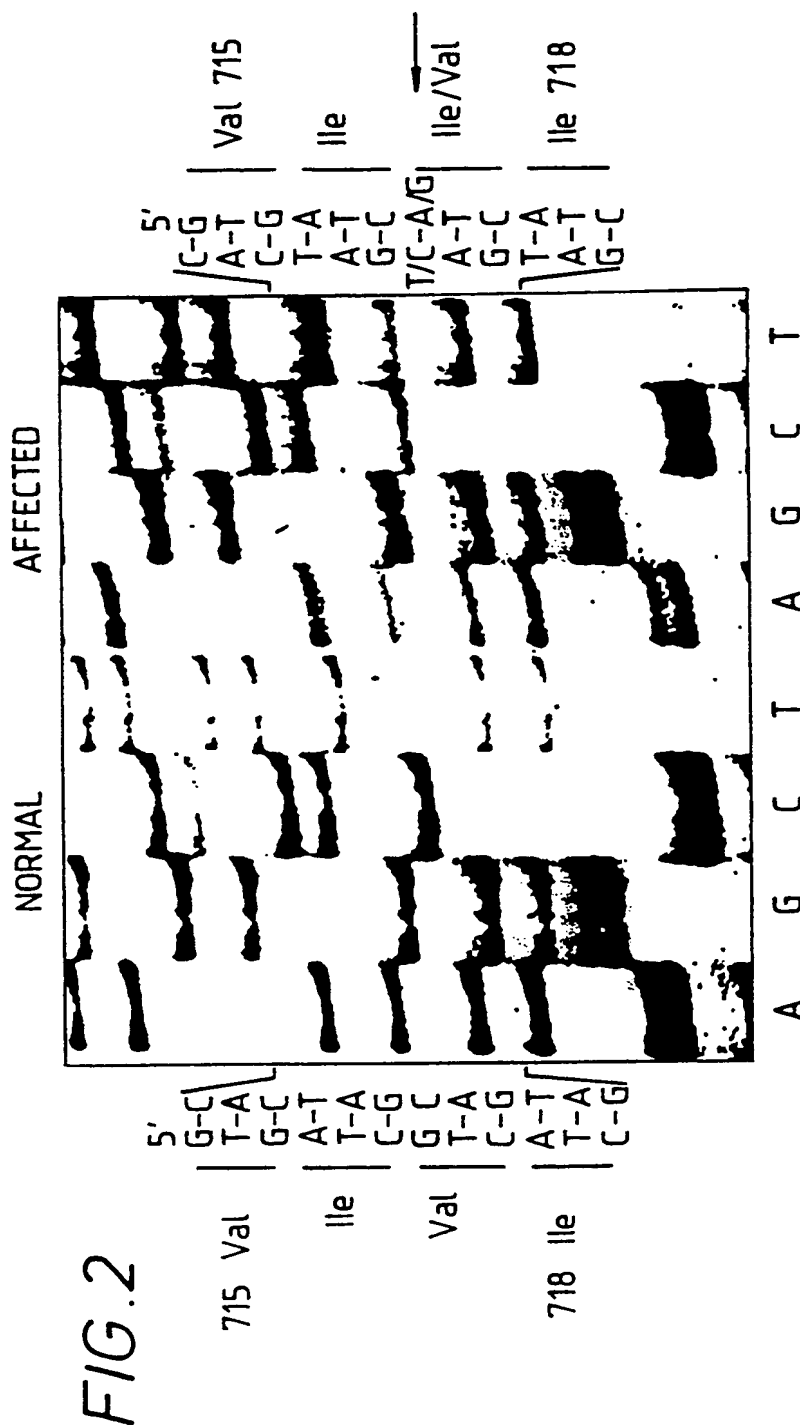
39. A method as claimed in claim 38, wherein a single amino acid is substituted for another.

5

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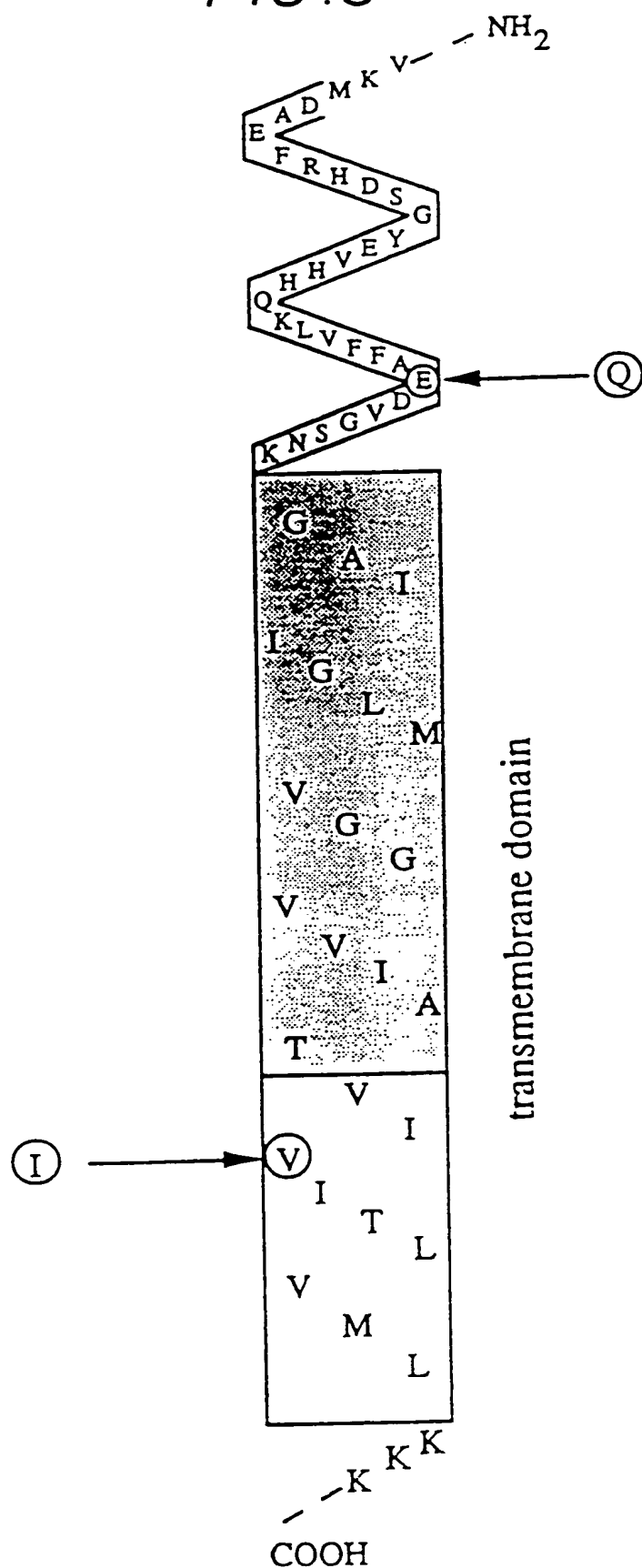


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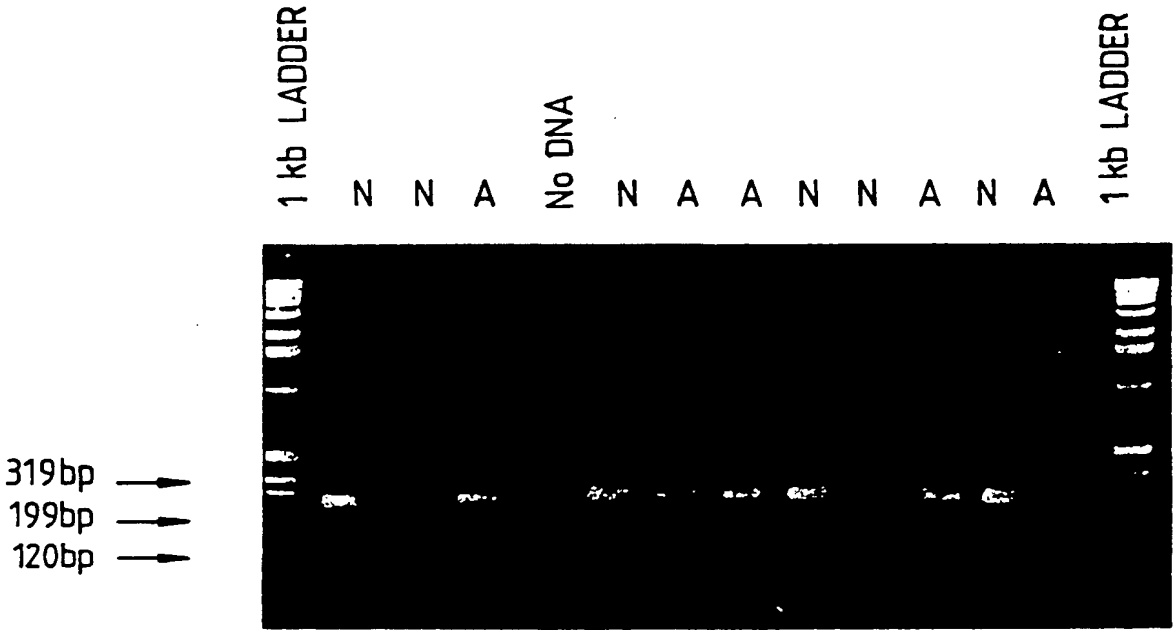
FIG. 3



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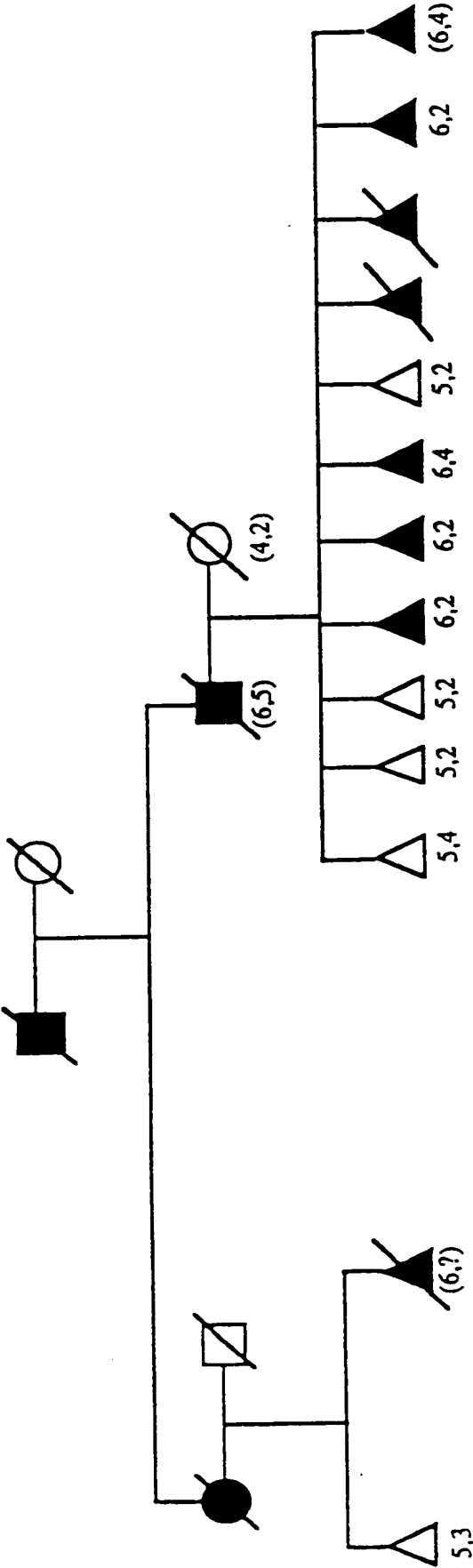
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FIG .4



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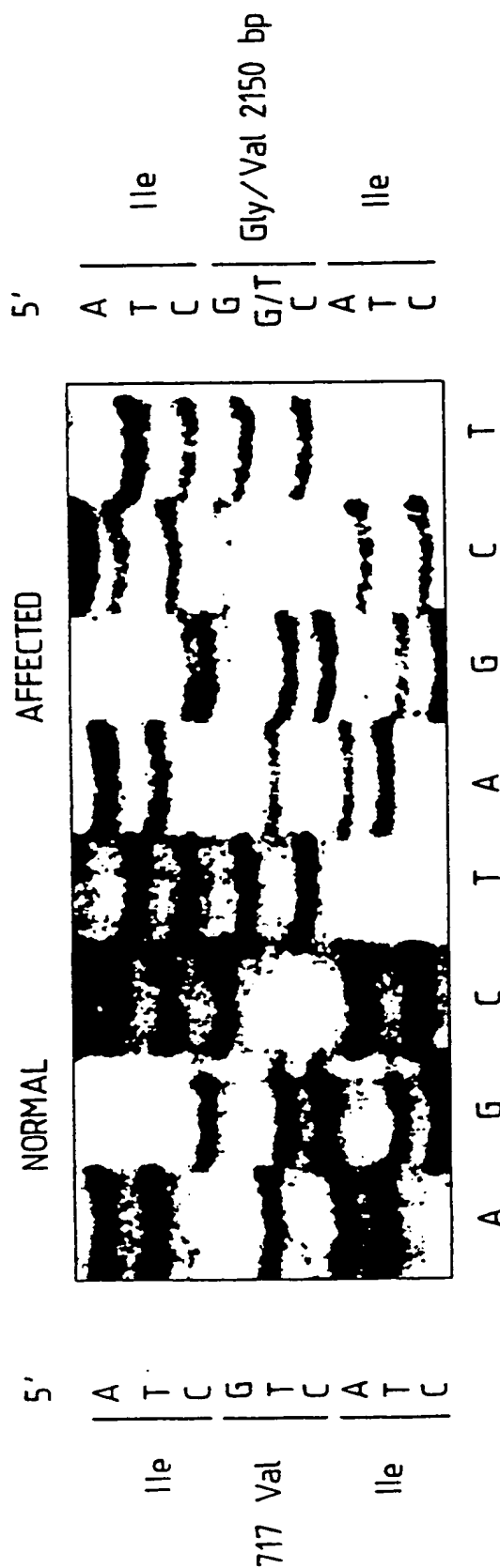
FIG. 5



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FIG. 6



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## FIG. 7 (1/7)

Met	Leu	Pro	Gly	Leu	Ala	Leu	Leu	Leu	Ala	Ala	Trp	Thr	Ala	Arg
1			5				10						15	
Ala	Leu	Glu	Val	Pro	Thr	Asp	Gly	Asn	Ala	Gly	Leu	Ala	Glu	Pro
		20					25					30		
Gln	Ile	Ala	Met	Phe	Cys	Gly	Arg	Leu	Asn	Met	His	Met	Asn	Val
		35					40					45		Gln
Asn	Gly	Lys	Trp	Asp	Ser	Asp	Pro	Ser	Gly	Thr	Lys	Thr	Cys	Ile
		50				55					60			Asp
Thr	Lys	Glu	Gly	Ile	Leu	Gln	Tyr	Cys	Gln	Glu	Val	Tyr	Pro	Glu
		65			70					75				80
Gln	Ile	Thr	Asn	Val	Val	Glu	Ala	Asn	Gln	Pro	Val	Thr	Ile	Gln
			85						90				95	Asn
Trp	Cys	Lys	Arg	Gly	Arg	Lys	Gln	Cys	Lys	Thr	His	Pro	His	Phe
			100					105					110	Val

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## FIG. 7 (217)

Ile	Pro	Tyr	Arg	Cys	Leu	Val	Gly	Glu	Phe	Val	Ser	Asp	Ala	Leu	Leu	
		115					120					125				
Val	Pro	Asp	Lys	Cys	Lys	Phe	Leu	His	Gln	Glu	Arg	Met	Asp	Val	Cys	
		130				135					140					
Glu	Thr	His	Leu	His	Trp	His	Thr	Val	Ala	Lys	Glu	Thr	Cys	Ser	Glu	
145					150					155					160	
Lys	Ser	Thr	Asn	Leu	His	Asp	Tyr	Gly	Met	Leu	Leu	Pro	Cys	Gly	Ile	
				165					170					175		
Asp	Lys	Phe	Arg	Gly	Val	Glu	Phe	Val	Cys	Cys	Pro	Leu	Ala	Glu	Glu	
			180					185					190			
Ser	Asp	Asn	Val	Asp	Ser	Ala	Asp	Ala	Glu	Glu	Asp	Asp	Ser	Asp	Val	
		195					200					205				

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## FIG. 7 (317)

Trp	Trp	Gly	Gly	Ala	Asp	Thr	Asp	Tyr	Ala	Asp	Gly	Ser	Glu	Asp	Lys
210						215					220				
	Val	Val	Glu	Val	Ala	Glu	Glu	Glu	Val	Ala	Glu	Val	Glu	Glu	Glu
225						230					235				240
	Glu	Ala	Asp	Asp	Glu	Asp	Asp	Glu	Asp	Gly	Asp	Glu	Val	Glu	Glu
						245					250				255
	Glu	Ala	Glu	Glu	Pro	Tyr	Glu	Glu	Ala	Thr	Glu	Arg	Thr	Ser	Ile
						260					265				270
	Ala	Thr	Thr	Thr	Thr	Thr	Thr	Thr	Glu	Ser	Val	Glu	Val	Val	Arg
						275									285
	Val	Pro	Thr	Thr	Ala	Ala	Ser	Thr	Pro	Asp	Ala	Val	Asp	Lys	Tyr
															Leu
	290						295					300			

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## FIG. 7 (4/7)

Glu Thr Pro Gly Asp	Glu Asn Glu His Ala His Phe Gln Lys Ala Lys	320
305	310	315
Glu Arg Leu Glu Ala	Lys His Arg Glu Arg Met Ser Gln Val Met Arg	335
325	330	
Glu Trp Glu Glu Ala Glu Arg Gln Ala Lys Asn Leu Pro Lys Ala Asp		350
340	345	
Lys Lys Ala Val Ile Gln His Phe Gln Glu Lys Val Glu Ser Leu Glu		365
355	360	
Gln Glu Ala Ala Asn Glu Arg Gln Gln Leu Val Glu Thr His Met Ala		380
370	375	
Arg Val Glu Ala Met Leu Asn Asp Arg Arg Arg Leu Ala Leu Glu Asn		400
385	390	395

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## FIG. 7 (5/7)

Tyr	Ile	Thr	Ala	Leu	Gln	Ala	Val	Pro	Pro	Arg	Pro	Arg	His	Val	Phe
				405					410					415	
Asn	Met	Leu	Lys	Lys	Tyr	Val	Arg	Ala	Glu	Gln	Lys	Asp	Arg	Gln	His
			420					425					430		
Thr	Leu	Lys	His	Phe	Glu	His	Val	Arg	Met	Val	Asp	Pro	Lys	Lys	Ala
		435					440					445			
Ala	Gln	Ile	Arg	Ser	Gln	Val	Met	Thr	His	Leu	Arg	Val	Ile	Tyr	Glu
	450					455					460				
Arg	Met	Asn	Gln	Ser	Leu	Ser	Leu	Leu	Tyr	Asn	Val	Pro	Ala	Val	Ala
	465				470					475					480
Glu	Glu	Ile	Gln	Asp	Glu	Val	Asp	Glu	Leu	Gln	Lys	Glu	Gln	Asn	
				485					490					495	

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## FIG. 7 (6/7)

Tyr Ser Asp Asp	Val Leu Ala Asn Met Ile Ser Glu Pro Arg Ile Ser
500	505 510
Tyr Gly Asn Asp	Pro Ser Leu Thr Glu Thr Lys Thr Thr
515	520 525
Val Glu Leu Leu Pro Val Asn Gly Glu Phe Ser Leu Asp Asp Leu Gln	
530	535 540
Pro Trp His Ser Phe Gly Ala Asp Ser Val Pro Ala Asn Thr Glu Asn	
545	550 555 560
Glu Val Glu Pro Val Asp Ala Arg Pro Ala Ala Asp Arg Gly Leu Thr	
565	570 575
Thr Arg Pro Gly Ser Gly Leu Thr Asn Ile Lys Thr Glu Glu Ile Ser	
580	585 590

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## FIG. 7 (7/7)

Glu Val Lys Met Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val	595	600	605
His His Gln Lys Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys	610	615	620
Gly Ala Ile Ile Gly Leu Met Val Gly Gly Val Val Ile Ala Thr Val	625	630	635
Ile Val Ile Thr Leu Val Met Leu Lys Lys Lys Gln Tyr Thr Ser Ile	645	650	655
His His Gly Val Val Glu Val Asp Ala Ala Val Thr Pro Glu Glu Arg	660	665	670
His Leu Ser Lys Met Gln Gln Asn Gly Tyr Glu Asn Pro Thr Tyr Lys	675	680	685
Phe Phe Glu Gln Met Gln Asn	690	695	

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## FIG. 8(1/8)

Met	Leu	Pro	Gly	Leu	Ala	Leu	Leu	Leu	Ala	Ala	Trp	Thr	Ala	Arg
1			5				10						15	
Ala	Leu	Glu	Val	Pro	Thr	Asp	Gly	Asn	Ala	Gly	Leu	Ala	Glu	Pro
		20					25					30		
Gln	Ile	Ala	Met	Phe	Cys	Gly	Arg	Leu	Asn	Met	His	Met	Asn	Val
		35					40					45		Gln
Asn	Gly	Lys	Trp	Asp	Ser	Asp	Pro	Ser	Gly	Thr	Lys	Thr	Cys	Ile
		50				55					60			Asp
Thr	Lys	Glu	Gly	Ile	Leu	Gln	Tyr	Cys	Gln	Glu	Val	Tyr	Pro	Glu
		65			70					75				80
Gln	Ile	Thr	Asn	Val	Val	Glu	Ala	Asn	Gln	Pro	Val	Thr	Ile	Gln
				85					90					95
Trp	Cys	Lys	Arg	Gly	Arg	Lys	Gln	Cys	Lys	Thr	His	Pro	His	Phe
			100						105					110

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## FIG. 8(2/8)

Ile	Pro	Tyr	Arg	Cys	Leu	Val	Gly	Glu	Phe	Val	Ser	Asp	Ala	Leu	Leu
		115					120					125			
Val	Pro	Asp	Lys	Cys	Lys	Phe	Leu	His	Gln	Glu	Arg	Met	Asp	Val	Cys
	130					135					140				
Glu	Thr	His	Leu	His	Trp	His	Thr	Val	Ala	Lys	Glu	Thr	Cys	Ser	Glu
145					150					155					160
Lys	Ser	Thr	Asn	Leu	His	Asp	Tyr	Gly	Met	Leu	Leu	Pro	Cys	Gly	Ile
				165				170						175	
Asp	Lys	Phe	Arg	Gly	Val	Glu	Phe	Val	Cys	Cys	Pro	Leu	Ala	Glu	Glu
			180					185					190		
Ser	Asp	Asn	Val	Asp	Ser	Ala	Asp	Ala	Glu	Glu	Asp	Asp	Ser	Asp	Val
		195					200					205			

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## FIG. 8(3/8)

Trp	Trp	Gly	Gly	Ala	Asp	Thr	Asp	Tyr	Ala	Asp	Gly	Ser	Glu	Asp	Lys
210						215					220				
	Val	Val	Glu	Val	Ala	Glu	Glu	Glu	Val	Ala	Glu	Val	Glu	Glu	Glu
225						230				235					240
	Glu	Ala	Asp	Asp	Glu	Asp	Asp	Glu	Asp	Gly	Asp	Glu	Val	Glu	Glu
					245				250						255
	Glu	Ala	Glu	Glu	Pro	Tyr	Glu	Glu	Ala	Thr	Glu	Arg	Thr	Ser	Ile
					260				265				270		
	Ala	Thr	Thr	Thr	Thr	Thr	Thr	Glu	Ser	Val	Glu	Glu	Val	Val	Arg
								280				285			
	Glu	Val	Cys	Ser	Glu	Gln	Ala	Glu	Thr	Gly	Pro	Cys	Arg	Ala	Met
290								295				300			

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## FIG. 8 (4/8)

Ser	Arg	Trp	Tyr	Phe	Asp	Val	Thr	Glu	Gly	Lys	Cys	Ala	Pro	Phe	Phe	320
305					310					315						
Tyr	Gly	Gly	Cys	Gly	Gly	Asn	Arg	Asn	Asn	Phe	Asp	Thr	Glu	Glu	Tyr	335
				325					330							
Cys	Met	Ala	Val	Cys	Gly	Ser	Ala	Ile	Pro	Thr	Thr	Ala	Ala	Ser	Thr	350
			340					345								
Pro	Asp	Ala	Val	Asp	Lys	Tyr	Leu	Glu	Thr	Pro	Gly	Asp	Glu	Asn	Glu	365
		355					360									
His	Ala	His	Phe	Gln	Lys	Ala	Lys	Glu	Arg	Leu	Glu	Ala	Lys	His	Arg	380
		370				375										
Glu	Arg	Met	Ser	Gln	Val	Met	Arg	Glu	Trp	Glu	Glu	Ala	Glu	Arg	Gln	400
385					390					395						

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## FIG. 8 (5/8)

Ala	Lys	Asn	Leu	Pro	Lys	Ala	Asp	Lys	Lys	Ala	Val	Ile	Gln	His	Phe
				405				410						415	
Gln	Glu	Lys	Val	Glu	Ser	Leu	Glu	Gln	Glu	Ala	Ala	Asn	Glu	Arg	Gln
			420					425					430		
Gln	Leu	Val	Glu	Thr	His	Met	Ala	Arg	Val	Glu	Ala	Met	Leu	Asn	Asp
		435					440					445			
Arg	Arg	Arg	Leu	Ala	Leu	Glu	Asn	Tyr	Ile	Thr	Ala	Leu	Gln	Ala	Val
		450				455					460				
Pro	Pro	Arg	Pro	Arg	His	Val	Phe	Asn	Met	Leu	Lys	Lys	Tyr	Val	Arg
					470					475					480
Ala	Glu	Gln	Lys	Asp	Arg	Gln	His	Thr	Leu	Lys	His	Phe	Glu	His	Val
				485					490					495	

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## FIG. 8 (6/8)

Arg Met Val Asp	Pro Lys Lys Ala Ala Gln Ile Arg Ser	Gln Val Met
500	505	510
Thr His Leu Arg Val Ile Tyr Glu Arg Met Asn Gln Ser	Leu Ser Leu	
515	520	525
Leu Tyr Asn Val Pro Ala Val Ala Glu Glu Ile Gln Asp Glu Val Asp		
530	535	540
Glu Leu Leu Gln Lys Glu Gln Asn Tyr Ser Asp Asp Val Leu Ala Asn		
545	550	555
Met Ile Ser Glu Pro Arg Ile Ser Tyr Gly Asn Asp Ala Leu Met Pro		
565	570	575
Ser Leu Thr Glu Thr Lys Thr Thr Val Glu Leu Leu Pro Val Asn Gly		
580	585	590

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FIG. 8 (7/8)

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## FIG. 8(8/8)

Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys Leu Val Phe Phe	660	665	670
Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile Gly Leu Met Val	675	680	685
Gly Gly Val Val Ile Ala Thr Val Ile Val Ile Thr Leu Val Met Leu	690	695	700
Lys Lys Lys Gln Tyr Thr Ser Ile His His Gly Val Val Glu Val Asp	705	710	715
Ala Ala Val Thr Pro Glu Glu Arg His Leu Ser Lys Met Gln Gln Asn	720	725	730
Gly Tyr Glu Asn Pro Thr Tyr Lys Phe Phe Glu Gln Met Gln Asn	735	740	745
			750



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## FIG. 9(1/8)

Met	Leu	Pro	Gly	Leu	Ala	Leu	Leu	Leu	Ala	Ala	Trp	Thr	Ala	Arg	1	5	10	15	
Ala	Leu	Glu	Val	Pro	Thr	Asp	Gly	Asn	Ala	Gly	Leu	Leu	Ala	Glu	Pro	20	25	30	
Gln	Ile	Ala	Met	Phe	Cys	Gly	Arg	Leu	Asn	Met	His	Met	Asn	Val	Gln	35	40	45	
Asn	Gly	Lys	Trp	Asp	Ser	Asp	Pro	Ser	Gly	Thr	Lys	Thr	Cys	Ile	Asp	50	55	60	
Thr	Lys	Glu	Gly	Ile	Leu	Gln	Tyr	Cys	Gln	Glu	Val	Tyr	Pro	Glu	Leu	65	70	75	80
Gln	Ile	Thr	Asn	Val	Val	Glu	Ala	Asn	Gln	Pro	Val	Thr	Ile	Gln	Asn	85	90	95	
Trp	Cys	Lys	Arg	Gly	Arg	Lys	Gln	Cys	Lys	Thr	His	Pro	His	Phe	Val	100	105	110	

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## FIG. 9 (2/8)

Ile	Pro	Tyr	Arg	Cys	Leu	Val	Gly	Glu	Phe	Val	Ser	Asp	Ala	Leu	Leu
		115					120					125			
Val	Pro	Asp	Lys	Cys	Lys	Phe	Leu	His	Gln	Glu	Arg	Met	Asp	Val	Cys
	130					135					140				
Glu	Thr	His	Leu	His	Trp	His	Thr	Val	Ala	Lys	Glu	Thr	Cys	Ser	Glu
	145				150					155					160
Lys	Ser	Thr	Asn	Leu	His	Asp	Tyr	Gly	Met	Leu	Leu	Pro	Cys	Gly	Ile
				165					170					175	
Asp	Lys	Phe	Arg	Gly	Val	Glu	Phe	Val	Cys	Cys	Pro	Leu	Ala	Glu	Glu
				180				185					190		
Ser	Asp	Asn	Val	Asp	Ser	Ala	Asp	Ala	Glu	Glu	Asp	Asp	Ser	Asp	Val
		195					200					205			

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## FIG. 9(3/8)

Trp	Trp	Gly	Gly	Ala	Asp	Thr	Asp	Tyr	Ala	Asp	Gly	Ser	Glu	Asp	Lys
210						215					220				
Val	Val	Glu	Val	Ala	Glu	Glu	Glu	Glu	Val	Ala	Glu	Val	Glu	Glu	Glu
225					230					235					240
Glu	Ala	Asp	Asp	Asp	Glu	Asp	Asp	Glu	Asp	Gly	Asp	Glu	Val	Glu	Glu
				245					250						255
Glu	Ala	Glu	Glu	Pro	Tyr	Glu	Glu	Ala	Thr	Glu	Arg	Thr	Thr	Ser	Ile
				260				265					270		
Ala	Thr	Thr	Thr	Thr	Thr	Thr	Thr	Glu	Ser	Val	Glu	Glu	Val	Val	Arg
							280					285			
Glu	Val	Cys	Ser	Glu	Gln	Ala	Glu	Thr	Gly	Pro	Cys	Arg	Ala	Met	Ile
290						295					300				

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## FIG. 9 (4/8)

Ser	Arg	Trp	Tyr	Phe	Asp	Val	Thr	Glu	Gly	Lys	Cys	Ala	Pro	Phe	Phe	320
305					310					315						
Tyr	Gly	Gly	Cys	Gly	Gly	Asn	Arg	Asn	Asn	Phe	Asp	Thr	Glu	Glu	Tyr	
									330						335	
Cys	Met	Ala	Val	Cys	Gly	Ser	Ala	Met	Ser	Gln	Ser	Leu	Leu	Lys	Thr	
									345						350	
Thr	Gln	Glu	Pro	Leu	Ala	Arg	Asp	Pro	Val	Lys	Leu	Pro	Thr	Thr	Ala	
Ala	Ser	Thr	Pro	Asp	Ala	Val	Asp	Lys	Tyr	Leu	Glu	Thr	Pro	Gly	Asp	
Glu	Asn	Glu	His	Ala	His	Phe	Gln	Lys	Ala	Lys	Glu	Arg	Leu	Glu	Ala	
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FIG. 9(5/8)

Lys	His	Arg	Glu	Arg	Met	Ser	Gln	Val	Met	Arg	Glu	Trp	Glu	Glu	Ala
				405					410					415	
Glu	Arg	Gln	Ala	Lys	Asn	Leu	Pro	Lys	Ala	Asp	Lys	Lys	Ala	Val	Ile
			420					425					430		
Gln	His	Phe	Gln	Glu	Lys	Val	Glu	Ser	Leu	Glu	Gln	Glu	Ala	Ala	Asn
		435					440					445			
Glu	Arg	Gln	Gln	Leu	Val	Glu	Thr	His	Met	Ala	Arg	Val	Glu	Ala	Met
	450						455				460				
Leu	Asn	Asp	Arg	Arg	Arg	Leu	Ala	Leu	Glu	Asn	Tyr	Ile	Thr	Ala	Leu
465					470					475					480
Gln	Ala	Val	Pro	Pro	Arg	Pro	Arg	His	Val	Phe	Asn	Met	Leu	Lys	Lys
					485				490					495	

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## FIG. 9(6/8)

Tyr	Val	Arg	Ala	Glu	Gln	Lys	Asp	Arg	Gln	His	Thr	Leu	Lys	His	Phe
			500					505					510		
Glu	His	Val	Arg	Met	Val	Asp	Pro	Lys	Lys	Ala	Ala	Gln	Ile	Arg	Ser
		515					520					525			
Gln	Val	Met	Thr	His	Leu	Arg	Val	Ile	Tyr	Glu	Arg	Met	Asn	Gln	Ser
		530				535					540				
Leu	Ser	Leu	Leu	Tyr	Asn	Val	Pro	Ala	Val	Ala	Glu	Glu	Ile	Gln	Asp
545					550					555					560
Glu	Val	Asp	Glu	Leu	Leu	Gln	Lys	Glu	Gln	Asn	Tyr	Ser	Asp	Asp	Val
				565					570					575	
Leu	Ala	Asn	Met	Ile	Ser	Glu	Pro	Arg	Ile	Ser	Tyr	Gly	Asn	Asp	Ala
			580					585					590		

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## FIG. 9(7/8)

Leu Met	Pro	Ser	Leu	Thr	Glu	Thr	Lys	Thr	Thr	Val	Glu	Leu	Leu	Pro
	595				600						605			
Val Asn	Gly	Glu	Phe	Ser	Leu	Asp	Asp	Leu	Gln	Pro	Trp	His	Ser	Phe
610					615					620				
Gly Ala	Asp	Ser	Val	Pro	Ala	Asn	Thr	Glu	Asn	Glu	Val	Glu	Pro	Val
625				630					635					640
Asp Ala	Arg	Pro	Ala	Ala	Asp	Arg	Gly	Leu	Thr	Thr	Arg	Pro	Gly	Ser
			645					650					655	
Gly Leu	Thr	Asn	Ile	Lys	Thr	Glu	Glu	Ile	Ser	Glu	Val	Lys	Met	Asp
							665					670		
Ala Glu	Phe	Arg	His	Asp	Ser	Gly	Tyr	Glu	Val	His	His	Gln	Lys	Leu
						680					685			
Val Phe	Phe	Ala	Glu	Asp	Val	Gly	Ser	Asn	Lys	Gly	Ala	Ile	Ile	Gly
690					695					700				

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FIG. 9 (8/8)

Val	Phe	Phe	Ala	Glu	Asp	Val	Gly	Ser	Asn	Lys	Gly	Ala	Ile	Ile	Gly
690						695					700				
Leu	Met	Val	Gly	Gly	Val	Val	Ile	Ala	Thr	Val	Ile	Val	Ile	Thr	Leu
705					710					715					720
Val	Met	Leu	Lys	Lys	Lys	Gln	Tyr	Thr	Ser	Ile	His	His	Gly	Val	Val
			725						730					735	
Glu	Val	Asp	Ala	Ala	Val	Thr	Pro	Glu	Glu	Arg	His	Leu	Ser	Lys	Met
			740			745							750		
Gln	Gln	Asn	Gly	Tyr	Glu	Asn	Pro	Thr	Tyr	Lys	Phe	Phe	Glu	Gln	Met
		755					760						765		
Gln	Asn														
770															

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